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OM nucleic - nucleic search, using sw model

Run on: October 31, 2002, 19:48:26 ; Search time 251 seconds

(without alignments)
13063.385 Million cell updates/sec

Title: US-09-919-197-3

Perfect score: 1456

Sequence: 1 tgagcaccgccaaccagg.....tatgaacactataaaaaa 1456

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 2185239 seqs, 112599159 residues

number of hits satisfying chosen parameters: 2331924

Minimum DB seq length: 0

Maximum DB seq length: 75

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : N_Geneseq_101002.*

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- 21: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2000.DAT.*
- 22: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2001A.DAT.*
- 23: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2001B.DAT.*
- 24: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2002.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	45	3.1	65	ABN52223	Mouse spliced tran
c 2	31.4	2.2	49	AAT89492	CFTR binding site
c 3	27.8	1.9	60	ABN49410	Human spliced tran
c 4	27.6	1.9	47	AA266146	Human map-related
5	26.2	1.8	47	AA265699	Human map-related
6	26	1.8	73	AA273834	Human HER2 partial
c 7	25.8	1.8	60	ABN40794	Human spliced tran
8	25.4	1.7	47	AAH38784	Human SNP flanking
c 9	25.4	1.7	60	ABN39098	Human spliced tran

c 10	25.4	1.7	72	19	AAT97276	Nucleotide EP-13 S
c 11	25.2	1.7	65	24	ABN28389	Rat spliced transc
c 12	25	1.7	51	22	AAL30738	Human SNP oligonuc
c 13	24.8	1.7	47	21	AA267790	Human map-related
c 14	24.6	1.7	60	24	ABN47816	Human spliced tran
c 15	24.4	1.7	51	22	AAL30269	Human SNP oligonuc
c 16	24	1.6	60	24	ABN39375	Human spliced tran
c 17	24	1.6	65	24	ABN28030	Rat spliced transc
c 18	23.8	1.6	64	21	AAAL1650	Humanised anti-Fas
c 19	23.8	1.6	64	24	ABL46029	Humanised anti-Fas
c 20	23.8	1.6	75	9	AAH80654	75-mer probe used
c 21	23.6	1.6	47	21	AAL266638	Human map-related
c 22	23.6	1.6	50	22	AAL31451	Human SNP oligonuc
c 23	23.6	1.6	60	18	AAT61582	VH and scFv antibo
c 24	23.6	1.6	60	24	ABN32752	Human spliced tran
c 25	23.6	1.6	60	24	ABN32853	Human spliced tran
c 26	23.6	1.6	60	24	ABN59247	Human spliced tran
c 27	23.6	1.6	71	18	AAT92616	SELEX-derived nucl
c 28	23.6	1.6	71	18	AAT78604	Class I SELEX gene
c 29	23.6	1.6	71	18	AAT59094	Human chorionic go
c 30	23	1.6	23	24	ABL45019	Human chromosome 1
c 31	23	1.6	51	22	AAL29070	Human SNP oligonuc
c 32	23	1.6	60	21	AAC23790	Human secreted pro
c 33	22.6	1.6	50	22	AAH89731	Human coding sequ
c 34	22.6	1.6	60	24	ABN32873	Human spliced tran
c 35	22.6	1.6	60	24	ABN34691	Human spliced tran
c 36	22.6	1.6	64	22	ABF80440	5-enolpyruvylshiki
c 37	22.6	1.6	69	15	AAQ44415	Special At-rich (A
c 38	22.4	1.5	50	22	AAH89839	Human coding sequ
c 39	22.4	1.5	60	24	ABN44442	Human spliced tran
c 40	22.4	1.5	68	22	AAL02305	Human reproductive
c 41	22.2	1.5	43	21	AAH75934	PCR primer used to
c 42	22.2	1.5	48	22	AAH22252	Anti-A33 antigen a
c 43	22.2	1.5	48	22	AAH20117	Rabbit anti A33 an
c 44	22.2	1.5	55	19	AAH70103	Humanised HFE7A he
c 45	22.2	1.5	55	20	AAH01319	Allelic ladder, HU

ALIGNMENTS

RESULT 1

ABN52223

ID ABN52223 standard; DNA; 65 BP.

XX AC ABN52223;

XX AC ABN52223;

DT 15-JUL-2002 (first entry)

XX Mouse spliced transcript detection oligonucleotide SEQ ID NO:24971.

DE Human; mouse; rat; splice transcript; detection; RNA transcript;

XX KW splice variant; transcriptome; oligonucleotide library; ss.

XX OS Mus musculus.

XX PN WO200210449-A2.

XX PD 07-FEB-2002.

XX 20-JUL-2001; 2001WO-IB01903.

XX 28-JUL-2000; 2000US-221607P.

XX 02-MAY-2001; 2001US-287724P.

XX (COMP-) COMPUGEN INC.

XX Shoshan A, Wasserman A, Mintz E, Mintz L, Faigler S;

XX WPI; 2002-257383/30.

XX New oligonucleotide libraries comprising oligonucleotides which

PT selectively hybridize to mRNAs transcribed from a transcription unit of

Fri NOV 1 18:25:06 2002

DR WPI; 1997-480211/44.
 XX New oligo:nucleotide(s) for gene therapy of cystic fibrosis - are
 PT able to increase the activity of the CF trans-membrane conductance
 PT regulator gene promoter
 XX
 PS Claim 3; Page 16; 20pp; English.
 XX
 CC This sequence represents a minimum sequence required for binding of
 CC proteins found in the nuclear extracts of cystic fibrosis transmembrane
 CC conductance regulator (CFTR)-expressing cells. This sequence is an
 CC oligonucleotide of the invention. This sequence, and sequences containing
 CC it increase the activity of the CFTR gene promoter, so regulate spatial
 CC expression of this gene (unlike viral or other promoters used in gene
 CC therapy). Constructs containing this sequence are used in gene therapy of
 CC cystic fibrosis (CF).
 XX
 CC Sequence 49 BP; 15 A; 10 C; 5 G; 19 T; 0 other;
 SQ
 Query Match 2.28; Score 31.4; DB 18; Length 49;
 Best Local Similarity 77.6%; Pred. No. 32;
 Matches 38; Conservative 0; Mismatches 11; Indels 0; Gaps 0;
 QY 557 AAGTCACTTGGCCCAAGGTCACCCAGCTAATAAGTCACAGTCTGGGATT 605
 DB 49 AAGTAAATTGCTTAGATCACATTGTTTAAAGTCACAGAGTAGGATT 1
 RESULT 3
 ID ABN49410/c
 XX ABN49410 standard; DNA; 60 BP.
 AC ABN49410;
 XX
 DT 15-JUL-2002 (first entry)
 XX
 DE Human spliced transcript detection oligonucleotide, SEQ ID NO:22158.
 XX
 KW Human; mouse; rat; splice transcript; detection; RNA transcript;
 KW splice variant; transcriptome; oligonucleotide library; ss.
 XX
 OS Homo sapiens.
 XX
 PN WO200210449-A2.
 XX
 PD 07-FEB-2002.
 XX
 PF 20-JUL-2001; 2001WO-1901903.
 XX
 PR 28-JUL-2000; 2000US-221607P.
 PR 02-MAY-2001; 2001US-287724P.
 XX
 PA (COMP-) COMPUGEN INC.
 XX
 PI Shoshan A, Wasserman A, Mintz E, Mintz L, Faigler S;
 XX
 XX WPI; 2002-257383/30.
 XX
 PT New oligonucleotide libraries comprising oligonucleotides which
 PT selectively hybridize to mRNAs transcribed from a transcription unit of
 PT a genome, useful for detecting tissue-, pathology-, and
 PT developmental-specific genes
 XX
 PS Example 1; SEQ ID 22158; 47pp; English.
 XX
 CC The present invention describes oligonucleotide libraries for detecting
 CC messenger RNAs that populate a (sub-)transcriptome, where the
 CC (sub-)transcriptome comprises messenger RNAs transcribed from multiple
 CC transcription units that populate a genome. The library comprises
 CC several oligonucleotides, each capable of hybridising selectively to a
 CC set of messenger RNAs transcribed from a given transcription unit of
 CC the genome, which encodes one or more messenger RNA splice variants.
 CC The oligonucleotide libraries are useful for detecting mRNAs from a
 CC biological sample, in expression profiling studies, in qualitatively or
 CC quantitatively characterising the corresponding transcriptome, and in
 CC detecting RNA transcripts and splice variants of human or animal
 CC transcriptomes. The libraries may also be used as specialised mini
 CC libraries to detect transcripts of a sub-transcriptome under a
 CC particular biological or pathological state, and so allowing the
 CC detection of tissue- and pathology-specific genes such as those genes
 CC only expressed in specific tissue under a specific pathological
 CC condition; to detect developmental specific genes; and to detect RNA
 CC transcripts and splice variants of a transcriptome of a patient suffering
 CC from a particular disorder. ABN27253 to ABN59589 represent
 CC oligonucleotide sequences from rats, humans and mice, which are used in
 CC the exemplification of the present invention.
 CC N.B. The sequence data for this patent did not form part of the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.
 XX
 SQ Sequence 65 BP; 15 A; 18 C; 14 G; 18 T; 0 other;

PT a genome, useful for detecting tissue-, pathology-, and
 XX developmental specific genes
 PT
 PS Example 1; SEQ ID 24971; 47pp; English.
 XX
 CC The present invention describes oligonucleotide libraries for detecting
 CC messenger RNAs that populate a (sub-)transcriptome, where the
 CC (sub-)transcriptome comprises messenger RNAs transcribed from multiple
 CC transcription units that populate a genome. The library comprises
 CC several oligonucleotides, each capable of hybridising selectively to a
 CC set of messenger RNAs transcribed from a given transcription unit of
 CC the genome, which encodes one or more messenger RNA splice variants.
 CC The oligonucleotide libraries are useful for detecting mRNAs from a
 CC biological sample, in expression profiling studies, in qualitatively or
 CC quantitatively characterising the corresponding transcriptome, and in
 CC detecting RNA transcripts and splice variants of human or animal
 CC transcriptomes. The libraries may also be used as specialised mini
 CC libraries to detect transcripts of a sub-transcriptome under a
 CC particular biological or pathological state, and so allowing the
 CC detection of tissue- and pathology-specific genes such as those genes
 CC only expressed in specific tissue under a specific pathological
 CC condition; to detect developmental specific genes; and to detect RNA
 CC transcripts and splice variants of a transcriptome of a patient suffering
 CC from a particular disorder. ABN27253 to ABN59589 represent
 CC oligonucleotide sequences from rats, humans and mice, which are used in
 CC the exemplification of the present invention.
 CC N.B. The sequence data for this patent did not form part of the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.
 XX
 SQ Sequence 65 BP; 15 A; 18 C; 14 G; 18 T; 0 other;
 Query Match 3.1%; Score 45; DB 24; Length 65;
 Best Local Similarity 83.6%; Pred. No. 0.004;
 Matches 51; Conservative 0; Mismatches 10; Indels 0; Gaps 0;
 QY 1017 GACCTCTCTTCGCTATCATTTGGAGATGTTGACATCGTGGCTTCTTTGGGACATG 1076
 DB 5 GATCTCTCTTCGCTATCATTTGGAGATGTTGACATCGTGGCTTCTTTGGGACATG 64
 QY 1077 C 1077
 DB 65 C 65
 RESULT 2
 AAT89492/c
 AAT89492 standard; DNA; 49 BP.
 AAT89492;
 XX
 DT 14-APR-1998 (first entry)
 XX
 DE CFTR binding site fragment 3.
 XX
 KW Cystic fibrosis transmembrane conductance regulator; CFTR; gene therapy;
 KW protein binding site; nuclear extract; gene promoter; cystic fibrosis;
 KW spatial expression regulator; CF; ss.
 XX
 OS Homo sapiens.
 XX
 PN WO9735005-A2.
 XX
 PD 25-SEP-1997.
 XX
 PF 20-MAR-1997; 97WO-GB00787.
 XX
 PR 20-MAR-1996; 96GB-0005808.
 XX
 PA (ISIS-) ISIS INNOVATION LTD.
 XX
 PI Harris A;
 XX

primers for the biallelic markers. The biallelic markers of the invention have a variety of uses: they can be used for high density mapping of the human genome, and in complex association studies and haplotyping studies which are useful in determining the genetic basis for disease states. Compositions and methods of the invention can also be useful for the identification of the targets for the development of pharmaceutical agents and diagnostic methods, as well as the characterisation of the differential efficacious responses to and side effects from pharmaceutical agents acting on a disease as well as other treatment.

N.B. The SEQ ID NOS 2852, 2913, 2974, 3035, 3096, 3157, 3227, 3297 and 3367, are not actually given a sequence in the Sequence Listing from the present invention.

Query Match : 1.98; Score 27.6; DB 21; Length 47;

```
Query Match      : 1.9%; Score 27.6; DB 21; Length 47;
Best Local Similarity 78.6%; Pred. No. 4e+02;
Matches 33; Conservative 0; Mismatches 9; Indels 0; Gaps 0;
```

QY 554 ATTAAGTGACTTGGCCCAAGGTCACCCAGCTAATAAGTGACAG 595
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Dd 47 ATTAAGTCACCTGGCCCAAGGTATCCCCCTATTAAAGTAGCTG 6

RESULT 5.

AAZ05099
ID AAZ65699 standard: DNA: 47 BP.

AA
AC
AA265699:

XX
XX

DT 11-SEP-2001 (first entry)

Human map-related biallelic marker SEO ID NO:46.

KW	Human genome; biallelic marker; high density disequilibrium map;
KW	genomic map; haplotype; phenotype; polymorphic base; genotyping;
KW	haplotyping; hybridisation; identification; characterisation;
KW	diagnosis; single nucleotide polymorphism; SNP; ds

OS Homo sapiens.

	Key	Location/Qualifiers replace(24,C)
FH		
FT	variation	

FT	standard_name=	"single nucleotide polymorphism"
11	/cay- a	

PN WO9954500-A2.

PD 28-OCT-1999.

21-APR-1999; 99WO-IB00822.

PR 21-APR-1998; 98US-0082614.

XX

XX
XX
XTCNED / TCTD)
WTXX
FT COLLEGE D, ВУЗЕНТЕТОXX
WFL; 2000-013261701.

PT Novel biallelic markers used to construct a high density disequilibrium
PT map of the human genome -

PS Claim 1; Page 240; 2745pp; English.

AAZ65654 to AAZ69578 represent human biallelic markers from the present invention, which contain a polymorphic base at position 24 of their nucleotide sequences. AAZ65979 to AAZ77440 represent amplification primers for the biallelic markers. The biallelic markers of the invention have a variety of uses: they can be used for high density mapping of the human genome, and in complex association studies and

CC only expressed in specific tissue under a specific pathological
CC condition; to detect developmental specific genes; and to detect RNA
CC transcripts and splice variants of a transcriptome of a patient suffering
CC from a particular disorder. ABN27253 to ABN59589 represent
CC oligonucleotide sequences from rats, humans and mice, which are used in
CC the exemplification of the present invention.
CC N.B. The sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.

XX
SQ Sequence 60 BP; 16 A; 15 C; 11 G; 18 T; 0 other;

Query Match 1.8%; Score 25.8; DB 24; Length 60;

Best Local Similarity 73.3%; Pred. No. 1.5e+03;

Matches 33; Conservative 0; Mismatches 12; Indels 0; Gaps 0;

OY 562 ACTTCCCAAGTCCACCCAGCTAATAAGTGACAGTCTGGGATTC 606

Db 60 ACTGGCTCTAGATCATGTAGCAATTAAGTGACAGAGCTGTGAATC 16

LT 8

AAH38784

ID AAH38784 standard; DNA; 47 BP.

XX AC AAH38784;

XX DT 14-AUG-2001 (first entry)

XX DE Human SNP flanking oligonucleotide SEQ ID 1580.

XX KW Single nucleotide polymorphism; SNP; single nucleotide primer extension;
XX SNPE; genotyping; agammaglobulinaemia; diabetes insipidus; cancer;
XX Lesch-Nyhan syndrome; muscular dystrophy; familial hypercholesterolaemia;
XX polycystic kidney disease; osteogenesis imperfecta; autoimmune disease;
XX acute intermittent porphyria; rheumatoid arthritis; multiple sclerosis;
XX inflammation; forensic investigation; paternity analysis; ds.

XX OS Homo sapiens.

XX PN WO200129262-A2.

XX PD 26-APR-2001.

XX PF 13-OCT-2000; 2000WO-US28436.

XX PR 15-OCT-1999; 99US-0160096.

XX (ORCH-) ORCHID BIOSCIENCES INC.

XX Picoult-Newburg L, Pohl M;

XX WPI; 2001-290930/30.

XX New genotyping oligonucleotide, useful for detecting the presence,
XX absence or identity of single polynucleotide polymorphism in a nucleic
XX acid sample

XX Claim 1; Page 58; 83pp; English.

XX Sequences AAH37205 - AAH40944 represent PCR primers, single nucleotide
XX primer extension (SNPE) primers, and the sequences of regions flanking
XX sites of single nucleotide polymorphisms SNPs. The present invention
XX includes kits for determining the presence or absence of a SNP, using the
XX oligonucleotides of the invention. The PCR primers are used to amplify a
XX SNP flanking sequence, the SNPE primer is used as a genotyping primer.
XX The oligonucleotides are useful for genotyping a nucleic acid sample by
XX performing a single-nucleotide primer extension reaction. The
XX oligonucleotides are useful for determining the presence, absence or
XX identity of a SNP and for genotyping nucleic acid samples, for e.g. to
XX assess by association analysis the genotype of an individual or group of
XX individuals, having a pathological phenotypic trait suspected of being
XX caused by one or more SNPs. Phenotypic traits include diseases e.g.

CC agammaglobulinaemia, diabetes insipidus, Lesch-Nyhan syndrome, muscular
CC dystrophy, familial hypercholesterolaemia, polycystic kidney disease,
CC osteogenesis imperfecta and acute intermittent porphyria. Phenotypic
CC traits also include symptoms of or susceptibility to multifactorial
CC disease of which a component is or may be genetic such as autoimmune
CC diseases, including, rheumatoid arthritis, multiple sclerosis,
CC inflammation, cancer, nervous system diseases and infection by pathogenic
CC microorganism. The method is also useful in forensic investigations and
CC paternity analysis. The present sequence represents a fragment of human
CC DNA flanking the site of a single nucleotide polymorphism.

XX SQ Sequence 47 BP; 12 A; 15 C; 11 G; 9 T; 0 other;

Query Match 1.7%; Score 25.4; DB 22;

Best Local Similarity 74.4%; Pred. No. 1.8e+03;

Matches 32; Conservative 0; Mismatches 11; Indels 0; Gaps 0;

OY 550 AGAGATTAAAGTACCTTGCCCAAGGTACCCAGCTAATAAGTGA 592

Db 4 AGAGGTCCAGCCACTTGCCCAAGGCCACACAGCTCTATTGCA 46

RESULT 9

ABN39098/c

ID ABN39098 standard; DNA; 60 BP.

XX AC ABN39098;

XX DT 15-JUL-2002 (first entry)

XX DE Human spliced transcript detection oligonucleotide SEQ ID NO:11846.

XX KW Human; mouse; rat; splice transcript; detection: RNA transcript;
XX splice variant; transcriptome; oligonucleotide library; ss.

XX OS Homo sapiens.

XX PN WO200210449-A2.

XX PD 07-FEB-2002.

XX PF 20-JUL-2001; 2001WO-IB01903.

XX PR 28-JUL-2000; 2000US-221607P.

XX PR 02-MAY-2001; 2001US-287724P.

XX PA (COMP-) COMPUGEN INC.

XX Shoshan A, Wasserman A, Mintz E, Mintz L, Faigler S;

XX WPI; 2002-257383/30.

XX New oligonucleotide libraries comprising oligonucleotides which
XX selectively hybridize to mRNAs transcribed from a transcription unit of
XX a genome, useful for detecting tissue-, pathology-, and
XX developmental-specific genes

XX Example 1; SEQ ID 11846; 47pp; English.

XX The present invention describes oligonucleotide libraries for detecting
XX messenger RNAs that populate a (sub-)transcriptome, where the
XX (sub-)transcriptome comprises messenger RNAs transcribed from multiple
XX transcription units that populate a genome. The library comprises
XX several oligonucleotides, each capable of hybridising selectively to a
XX set of messenger RNAs transcribed from a given transcription unit of
XX the genome, which encodes one or more messenger RNA splice variants.
XX The oligonucleotide libraries are useful for detecting mRNAs from a
XX biological sample, in expression profiling studies, in qualitatively or
XX quantitatively characterising the corresponding transcriptome, and in
XX detecting RNA transcripts and splice variants of human or animal
XX transcriptomes. The libraries may also be used as specialised mini
XX libraries to detect transcripts of a sub-transcriptome under a
XX particular biological or pathological state, and so allowing the

detection of tissue- and pathology-specific genes such as those genes only expressed in specific tissue under a specific pathological condition; to detect developmental specific genes; and to detect RNA transcripts and splice variants of a transcriptome of a patient suffering from a particular disorder. ABN27253 to ABN59589 represent oligonucleotide sequences from rats, humans and mice, which are used in the exemplification of the present invention.

N.B. The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from WIPO at www.int/pub/published/pct_sequences.

xx
SQ sequence 60 BP; 14 A; 16 C; 15 G; 15 T; 0 other;
Query Match 1.7%; Score 25.4; DB 24; Length 60;
Best Local Similarity 74.4%; Pred. No. 2e+03;
Matches 32; Conservative 0; Mismatches 11; Indels 0; Gaps 0;

QY
562 ACTTGCCCAAGGTACCCAGCTAATAAGTGACAGTCTGGGAT 604
|||||
56 ACTTTGCTGAGGTACACTTGTCTGAAGTGACACAGCTGGCAT 14

RESULT 10
AAT97276/C
ID: AAT97276 standard; DNA: 72 BP.

AC AAT97276;
XX
DT 11-MAR-1998 (first entry)

XX Nucleotide Ep-13 SEQ ID NO:51 from J09278794 Example 7.
DE

XX Hepatitis C virus; HCV; chimeric; antigen; detection; core region;
KW NS3; NS4; infection; PCR primer; ss.

XX	Synthetic.	JP09278794-A
OS		
XX		
PN		

XX 28-OCT-1997 XX 0375-0027015

XX 09-FEB-1996; 96JP-0024045.

XX PA (TOFU) TONEN CORP.

XX
WPT. 1009-022218/03

New chimeric peptide antigen derived from hepatitis C virus protein - useful for detecting HCV infections

XX
PS Example 7: Page 14; 30pp; Japanese.

xx CC The present sequence represents a nucleotide sequence from example 7
CC of the present invention describing a HCV chimeric peptide antigen
CC which comprises at least 2 peptide epitope regions from the HCV
CC polypeptide core region, 2 peptide epitope regions from the NS3 region
CC and at least 2 peptide epitope regions from the NS4 region. The antigen
CC binds specifically with an antibody produced by a human infected by
CC HCV. The peptide can detect a wide range of HCV infections with high
CC sensitivity.

xx
sequence 72 BP; 15 A; 25 C; 21 G; 11 T; 0 other;

Query Match	1.7%	Score 25.4;	DB 19;	Length 72;
Best Local Similarity	68.6%	Pred. N.M. 2.2e+03;		
Matches	35;	Conservative	0;	Indels 0;
		Matches	16;	Gaps 0;

QY 1081 TGCTCAGGTGACCTGTTCCAGCCCCAGGCAGAGATCAGGTGGCAGAGGCTG 1131

70 TGGTGGCGCTACCGTAAATACCTGGGGACGGCGGTG 20

RESULT 11	
ABN28389/C	
ID	ABN28389 standard; DNA; 65 BP.
XX	
XX	
AC	ABN28389;
XX	
DT	15-JUL-2002 (first entry)
XX	
XX	rat spliced transcript detection; oligonucleotide SEQ ID NO:1137.
DE	

XX. Human; mouse; rat; splice transcript; detection; RNA transcript;
KW splice variant; transcriptome; oligonucleotide library; ss.

XX OS Rattus norvegicus.

XX
PN
W0200210449-A2

XX
XX

XX

XX
FF

PR 28-JUL-2000; 2000US 2210071
PR 02-MAY-2001; 2001US-287724P.

[illegible]

XX

XX
XX

DR WPL; 2002-231303/30;
XX

pt. New oligonucleotide
pt. selectively hybridiz-

PT a genome, useful for

XX.

XX
12
Lump

the present invention comprises messenger RNAs transcribed from multiple (sub-)transcriptome units that populate a genome. The library comprises several oligonucleotides, each capable of hybridising selectively to a given transcription unit of a (sub-)transcriptome, where the messenger RNAs that populate a (sub-)transcriptome, where the

CC set of messenger RNAs transcribed from the
CC genome, which encodes one or more messenger RNA splice variants.
CC The oligonucleotide libraries are useful for detecting mRNAs from a
CC biological sample, in expression profiling studies, in qualitatively
CC quantitatively characterising the corresponding transcriptome, and in
CC quantitatively characterising the splice variants of human or animal

CC detecting RNA transcripts and splice variants of named genes
CC transcriptomes. The libraries may also be used as specialised mini
CC libraries to detect transcripts of a sub-transcriptome under a
CC particular biological or pathological state, and so allowing the
CC detection of tissue- and pathology-specific genes such as those genes
CC

only expressed in specific tissue under a specific pathologic condition; and to detect RNA condition; to detect developmental specific genes; and to detect RNA transcripts and splice variants of a transcriptome of a patient suffering from a particular disorder. ABN27253 to ABN59589 represent oligonucleotide sequences from rats, humans and mice, which are used in

CC the exemplification of the present invention.
CC N.B. The sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.

XX sequence 65 BP; 8 A; 23 C; 14 G; 20 T; 0 other; 50

Query Match	1.78;	Score	25.2;	DB 24;	Length	65;
Query Similarity	71.78;	Bred NO	2.4e+03;			

Best Local Similarity	21.7%	Indels	0;
Matches	33;	Mismatches	-13;
	Conservative		Gaps

707 ACCACCACTAGTGGACCTCAAGGCCGAGCAAGGAGGCAG 752

18. CAGGGCCTCCCTCCAGGAGGCAG

10

RESULT 12

AAAL30738/c
ID AAL30738 standard; DNA; 51 BP.

XX
AC AAL30738;
XX

DT 24-JAN-2002 (first entry)
XX

DE Human SNP oligonucleotide #3946.
XX

XX Immunosuppressive; immunostimulatory; antiinflammatory; cytostatic;
KW neuroprotective; antimicrobial; gene therapy; vaccine; amylose; cancer;
KW amyloid protein; angiotensin; apoptosis related protein; cadherin;
KW cyclin; polymerase; oncogene; histone; kinase; colony stimulating factor;
KW complement related protein; cytochrome; kinesin; cytokine; interferon;
KW interleukin; G-protein coupled receptor; thioesterase; inflammation;
KW multifactorial disease; autoimmune disease; infection;
KW nervous system disease; ss.

XX Homo sapiens.

PN WO200147944-A2.

PD 05-JUL-2001.

PF 28-DEC-2000; 2000WO-US35498.

PR 28-DEC-1999; 99US-0173419.

PR 27-DEC-2000; 2000US-0173419.

XX (CURA-) CURAGEN CORP.

PA Shimkets RA, Leach M;

PI WPT; 2001-465210/50.

DR Polymorphic nucleic acids encoding e.g. amylases, cyclins, polymerases,
XX oncogenes and histones, useful for diagnosing and treating, e.g.
PT cancer, autoimmune diseases and infections

XX Claim 1; Page 2520; 4143pp; English.

XX The present invention relates to oligonucleotides encoding polymorphic
CC variants of proteins related to amylases, amyloid proteins, angiotensin,
CC apoptosis related proteins, cadherin, cyclin, polymerase, oncogenes,
CC histones, kinases, colony stimulating factors, complement related
CC proteins, cytochromes, kinesins, cytokines, interferons, interleukins,
CC G-protein coupled receptors and thioesterases. The present sequence is
CC one such oligonucleotide. The oligonucleotides and the peptides encoded
CC by them may be used in the prevention, diagnosis and treatment of
CC diseases associated with inappropriate expression of the proteins listed
CC above. Disorders that may be prevented, diagnosed and/or treated include
CC multifactorial diseases with a genetic component, such as autoimmune
CC diseases (e.g. rheumatoid arthritis, multiple sclerosis, diabetes,
CC systemic lupus erythematosus and Grave's disease), inflammation, cancer
CC (e.g. cancers of the bladder, brain, breast, colon and kidney,
CC leukaemia), diseases of the nervous system and an infection of pathogenic
CC organisms.

XX Sequence 51 BP; 13 A; 5 C; 1 G; 32 T; 0 other;

Query Match 1.7%; Score 25; DB 22; Length 51;
Best Local Similarity 69.4%; Pred. No. 2.4e+03;
Matches 34; Conservative 0; Mismatches 15; Indels 0; Gaps 0;

QY 1408 AATAAGTATTGATTTTGTATTAATAAGGTATGAACACTAAAAA 1456

DB 49 ACTAAAGTGCTATTATTTGTAGTAAAAA 1

RESULT 13

AAZ67790

AAZ67790 standard; DNA; 47 BP.

XX AAZ67790;

DT 10-SEP-2001 (first entry)
XX

DE Human map-related biallelic marker SEQ ID NO:2137.

XX Human genome; biallelic marker; high density disequilibrium map;
KW genomic map; haplotype; phenotype; polymorphic base; genotyping;
KW haplotyping; hybridisation; identification; characterisation;
KW diagnosis; single nucleotide polymorphism; SNP; ds.

XX Homo sapiens.

XX Key Location/Qualifiers

FT variation replace(24,G)

FT /*tag= a

FT /standard_name= "single nucleotide polymorphism"

PN WO9954500-A2.

PD 28-OCT-1999.

PF 21-APR-1999; 99WO-IB00822.

PR 21-APR-1998; 98US-0082614.

PR 23-NOV-1998; 98US-0109732.

XX (GEST) GENSET.

XX Cohen D, Blumenfeld M, Chumakov I;

PI WPI; 2000-013267/01.

XX Novel biallelic markers used to construct a high density disequilibrium
PT map of the human genome

PS Claim 1; Page 686; 2745pp; English.

XX AAZ65654 to AAZ69578 represent human biallelic markers from the present
CC invention, which contain a polymorphic base at position 24 of their
CC nucleotide sequences. AAZ69579 to AAZ77440 represent amplification
CC primers for the biallelic markers. The biallelic markers of the
CC invention have a variety of uses: they can be used for high density
CC mapping of the human genome, and in complex association studies and
CC haplotyping studies which are useful in determining the genetic basis
CC for disease states. Compositions and methods of the invention can also
CC be useful for the identification of the targets for the development of
CC pharmaceutical agents and diagnostic methods, as well as the
CC characterisation of the differential efficacious responses to and side
CC effects from pharmaceutical agents acting on a disease as well as other
CC treatment.

CC N.B. the SEQ ID NOS 2852, 2913, 2974, 3035, 3096, 3157, 3227, 3297
CC and 3367, are not actually given a sequence in the Sequence Listing
CC from the present invention.

XX Sequence 47 BP; 16 A; 10 C; 11 G; 10 T; 0 other;

Query Match 1.7%; Score 24.8; DB 21; Length 47;
Best Local Similarity 72.7%; Pred. No. 2.6e+03;
Matches 32; Conservative 0; Mismatches 13; Indels 0; Gaps 0;

QY 550 AGAGATTAAAGTACCTGCCCAAGGTACCCACCTCAATTAAGTAC 593

DB 3 AGAAACAAGGACACTTGGCCCAAGGTGCTGCAATTAAGTACCC 46

RESULT 14

ABN47816

ID ABN47816 standard; DNA; 60 BP.

XX ABN47816;

XX DT 15-JUL-2002 (first entry)
 XX DE Human spliced transcript-detection oligonucleotide SEQ ID NO:20564.
 XX KW Human; mouse; rat; splice transcript; detection; RNA transcript;
 XX KW splice variant; transcriptome; oligonucleotide library; ss.
 XX OS Homo sapiens
 XX PN WO200210449-A2.
 XX PD 07-FEB-2002.
 XX PF 20-JUL-2001; 2001WO-IB01903.
 XX PR 28-JUL-2000; 2000US-221607P.
 XX PR 02-MAY-2001; 2001US-287724P.
 XX (COMP-) COMPUGEN INC.
 XX Shoshan A, Wasserman A, Mintz E, Mintz L, Faigler S;
 XX WPI; 2002-257383/30..
 XX New oligonucleotide libraries comprising oligonucleotides which
 XX selectively hybridize to mRNAs transcribed from a transcription unit of
 XX a genome, useful for detecting tissue-, pathology-, and
 XX developmental-specific genes
 XX Example 1; SEQ ID 20564; 47pp; English.
 XX The present invention describes oligonucleotide libraries for detecting
 XX messenger RNAs that populate a (sub-)transcriptome, where the
 XX (sub-)transcriptome comprises messenger RNAs transcribed from multiple
 XX transcription units that populate a genome. The library comprises
 XX several oligonucleotides, each capable of hybridizing selectively to a
 XX set of messenger RNAs transcribed from a given transcription unit of
 XX the genome, which encodes one or more messenger RNA splice variants.
 XX The oligonucleotide libraries are useful for detecting mRNAs from a
 XX biological sample, in expression profiling studies, in qualitatively or
 XX quantitatively characterizing the corresponding transcriptome, and in
 XX detecting RNA transcripts and splice variants of human or animal
 XX transcriptomes. The libraries may also be used as specialised mini
 XX libraries to detect transcripts of a sub-transcriptome under a
 XX particular biological or pathological state, and so allowing the
 XX detection of tissue- and pathology-specific genes such as those genes
 XX only expressed in specific tissue under a specific pathological
 XX condition; to detect developmental specific genes; and to detect RNA
 XX transcripts and splice variants of a transcriptome of a patient suffering
 XX from a particular disorder. ABN27253 to ABN59589 represent
 XX oligonucleotide sequences from rats, humans and mice, which are used in
 XX the exemplification of the present invention.
 XX N.B. The sequence data for this patent did not form part of the printed
 XX specification, but was obtained in electronic format directly from WIPO
 XX at ftp.wipo.int/pub/published_pct_sequences.
 XX Sequence 60 BP; 20 A; 13 C; 15 G; 12 T; 0 other;
 XX Query Match 1.7%; Score 24.6; DB 24; Length 60;
 XX Best Local Similarity 65.5%; Pred. No. 3.5e+03;
 XX Matches 36; Conservative 0; Mismatches 19; Indels 0; Gaps 0;
 QY 547 CCAGAGATTAGTACTTCCACAGCTCAGCTCAATAGTGACAGTGCTGG 601
 DB 1 CCTAGAGTTAGTTACTTCTGCTCAAGACACAGTAGCTAGTAGGACCAAGCCAGG 55
 RESULT 15
 AAL30269/C
 ID AAL30269 standard; DNA; 51 BP.
 XX AAL30269;
 XX

XX DT 24-JAN-2002 (first entry)
 XX DE Human SNP oligonucleotide #3477.
 XX KW Immunosuppressive; immunostimulatory; antiinflammatory; cytostatic;
 XX KW neuroprotective; antimicrobial; gene therapy; vaccine; amylase; cancer;
 XX KW amyloid protein; angiotensin; apoptosis related protein; cadherin;
 XX KW cyclin; polymerase; oncogene; histone; kinase; colony stimulating factor;
 XX KW complement related protein; cytochrome; kinase; cytokine; interferon;
 XX KW interleukin; G-protein coupled receptor; thioesterase; inflammation;
 XX KW multifactorial disease; autoimmune disease; infection;
 XX KW nervous system disease; ss.
 XX OS Homo sapiens.
 XX PN WO200147944-A2.
 XX PD 05-JUL-2001.
 XX PF 28-DEC-2000; 2000WO-US35498.
 XX PR 28-DEC-1999; 99US-0173419.
 XX PR 27-DEC-2000; 2000US-0173419.
 XX (CURA-) CURAGEN CORP.
 XX Shimkets RA, Leach M;
 XX WPI; 2001-465210/50.
 XX Polymorphic nucleic acids encoding e.g. amylases, cyclins, polymerases,
 XX oncogenes and histones, useful for diagnosing and treating, e.g.
 XX cancer, autoimmune diseases and infections
 XX Claim 1; Page 2383; 4143pp; English.
 XX The present invention relates to oligonucleotides encoding polymorphic
 XX variants of proteins related to amylases, amyloid proteins, angiotensin,
 XX apoptosis related proteins, cadherin, cyclin, polymerase, oncogenes,
 XX histones, kinases, colony stimulating factors, complement related
 XX proteins, cytochromes, kinesins, cytokines, interferons, interleukins,
 XX G-protein coupled receptors and thioesterases. The present sequence is
 XX one such oligonucleotide. The oligonucleotides and the peptides encoded
 XX by them may be used in the prevention, diagnosis and treatment of
 XX diseases associated with inappropriate expression of the proteins listed
 XX above. Disorders that may be prevented, diagnosed and/or treated include
 XX multifactorial diseases with a genetic component, such as autoimmune
 XX diseases (e.g. rheumatoid arthritis, multiple sclerosis, diabetes,
 XX systemic lupus erythematosus and Grave's disease), inflammation, cancer
 XX (e.g. cancers of the bladder, brain, breast, colon and kidney,
 XX leukaemia), diseases of the nervous system and an infection of pathogenic
 XX organisms.
 XX Sequence 51 BP; 12 A; 1 C; 2 G; 36 T; 0 other;
 XX Query Match 1.7%; Score 24.4; DB 22; Length 51;
 XX Best Local Similarity 68.0%; Pred. No. 3.6e+03;
 XX Matches 34; Conservative 0; Mismatches 16; Indels 0; Gaps 0;
 QY 1407 AATTAAGTTATTGATTTTGTAAATAAAGGTATGAACACATAAAAAA 1456
 DB 51 AATAAAATCTTATCTTTAGTAAAAAATAAAAAAATAAAAAAATAAAAAA 2
 Search completed: October 31, 2002, 21:55:33
 Job time : 253 secs

GenCore version 5.1.3
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OM nucleic - nucleic search, using sw model

Run on: October 31, 2002, 19:54:31 ; Search time 2921 Seconds
(without alignments)
14506.568 Million cell updates/sec

Title: US-09-919-197-3
Perfect score: 1456
Sequence: 1 tgaacaccagccaccagg.....tatgaacacataaaaaaa 1456

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues

1 number of hits satisfying chosen parameters: 941340

Minimum DB seq length: 0
Maximum DB seq length: 75

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : GenEmbl.*

- 1: gb_ba.*
- 2: gb_htg.*
- 3: gb_in.*
- 4: gb_on.*
- 5: gb_ov.*
- 6: gb_pat.*
- 7: gb_ph.*
- 8: gb_pi.*
- 9: gb_pr.*
- 10: gb_ro.*
- 11: gb_sts.*
- 12: gb_sy.*
- 13: gb_un.*
- 14: gb_vl.*
- 15: em_ba.*
- 16: em_fun.*
- 17: em_hum.*
- 18: em_in.*
- 19: em_mu.*
- 20: em_or.*
- 21: em_ov.*
- 22: em_pat.*
- 23: em_ph.*
- 24: em_pi.*
- 25: em_ro.*
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- 27: em_un.*
- 28: em_vl.*
- 29: em_vr.*
- 30: em_htg_hum.*
- 31: em_htg_inv.*
- 32: em_htg_other.*
- 33: em_htg_mus.*
- 34: em_htg_pln.*
- 35: em_htg_rtd.*
- 36: em_htg_mam.*
- 37: em_htg_vrt.*
- 38: em_sy.*
- 39: em_htgo_hum.*
- 40: em_htgo_mus.*
- 41: em_htgo_other.*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB	ID	Description
C 1	31.4	2.2	49	6	A65431	A65431 Sequence 23
C 2	29.4	2.0	71	4	AF055530	AF055530 Didelphis
C 3	25.4	1.7	47	6	AX116457	AX116457 Sequence
C 4	25.4	1.7	72	6	E14197	E14197 PCR primer
C 5	24.6	1.7	64	9	HS001823	AJ001823 Homo sapi
C 6	23.8	1.6	64	6	E40916	E40916 Humanized a
C 7	23.8	1.6	75	6	A05173	A05173 Oligonucleo
C 8	23.6	1.6	60	6	AR117953	AR117953 Sequence
C 9	23.6	1.6	71	6	AR020462	AR020462 Sequence
C 10	23.6	1.6	71	6	AR054774	AR054774 Sequence
C 11	23.6	1.6	71	6	AR066039	AR066039 Sequence
C 12	23.4	1.6	75	6	A15067	A15067 Nucleotide
C 13	23	1.6	74	6	A08223	A08223 synthetic o
C 14	23	1.6	74	6	A08224	A08224 synthetic o
C 15	22.8	1.6	74	8	CBE345111	AJ345111 Carpinus
C 16	22.6	1.6	50	6	AX199582	AX199582 Sequence
C 17	22.6	1.6	65	6	AX482787	AX482787 Sequence
C 18	22.4	1.5	50	6	AX199690	AX199690 Sequence
C 19	22.4	1.5	54	6	AR139994	AR139994 Sequence
C 20	22.4	1.5	65	6	AX485185	AX485185 Sequence
C 21	22.2	1.5	48	6	AR184251	AR184251 Sequence
C 22	22.2	1.5	48	6	AR184372	AR184372 Sequence
C 23	22.2	1.5	48	6	AX127963	AX127963 Sequence
C 24	22.2	1.5	55	6	AX002002	AX002002 Sequence
C 25	22.2	1.5	55	6	E40050	E40050 Drug contai
C 26	22.2	1.5	55	6	E40858	E40858 Humanized a
C 27	22.2	1.5	55	6	E43404	E43404 Humanized a
C 28	22.2	1.5	61	6	AR051730	AR051730 Sequence
C 29	22.2	1.5	61	6	AR117040	AR117040 Sequence
C 30	22.2	1.5	61	6	AR200053	AR200053 Sequence
C 31	22.2	1.5	61	6	I20500	I20500 Sequence 22
C 32	22.2	1.5	61	6	I49676	I49676 Sequence 20
C 33	22.2	1.5	63	6	BD002020	BD002020 Construct
C 34	22.2	1.5	63	6	I08751	I08751 Sequence 13
C 35	22.2	1.5	65	6	AR117041	AR117041 Sequence
C 36	22.2	1.5	65	6	AR200054	AR200054 Sequence
C 37	22.2	1.5	65	6	I20501	I20501 Sequence 23
C 38	22.2	1.5	65	6	I49677	I49677 Sequence 21
C 39	22.2	1.5	69	6	I16689	I16689 Sequence 29
C 40	22.2	1.5	69	6	I89442	I89442 Sequence 29
C 41	22.2	1.5	70	6	AX328443	AX328443 Sequence
C 42	22	1.5	39	6	AX137540	AX137540 Sequence
C 43	22	1.5	51	6	AX162353	AX162353 Sequence
C 44	22	1.5	58	6	E03095	E03095 RNA encodin
C 45	22	1.5	59	9	S76156	S76156 delta-beta

ALIGNMENTS

RESULT 1
A65431/c
LOCUS A65431
DEFINITION Sequence 23 from Patent WO9735005.
ACCESSION A65431
VERSION A65431.1 GI:4531190
KEYWORDS
SOURCE unidentified.
ORGANISM unidentified.
REFERENCE 1 (bases 1 to 49)
AUTHORS Harris A.
TITLE CFTR GENE REGULATOR
JOURNAL Patent: WO 9735005-A 23 25-SEP-1997;
ISIS INNOVATION (GB)

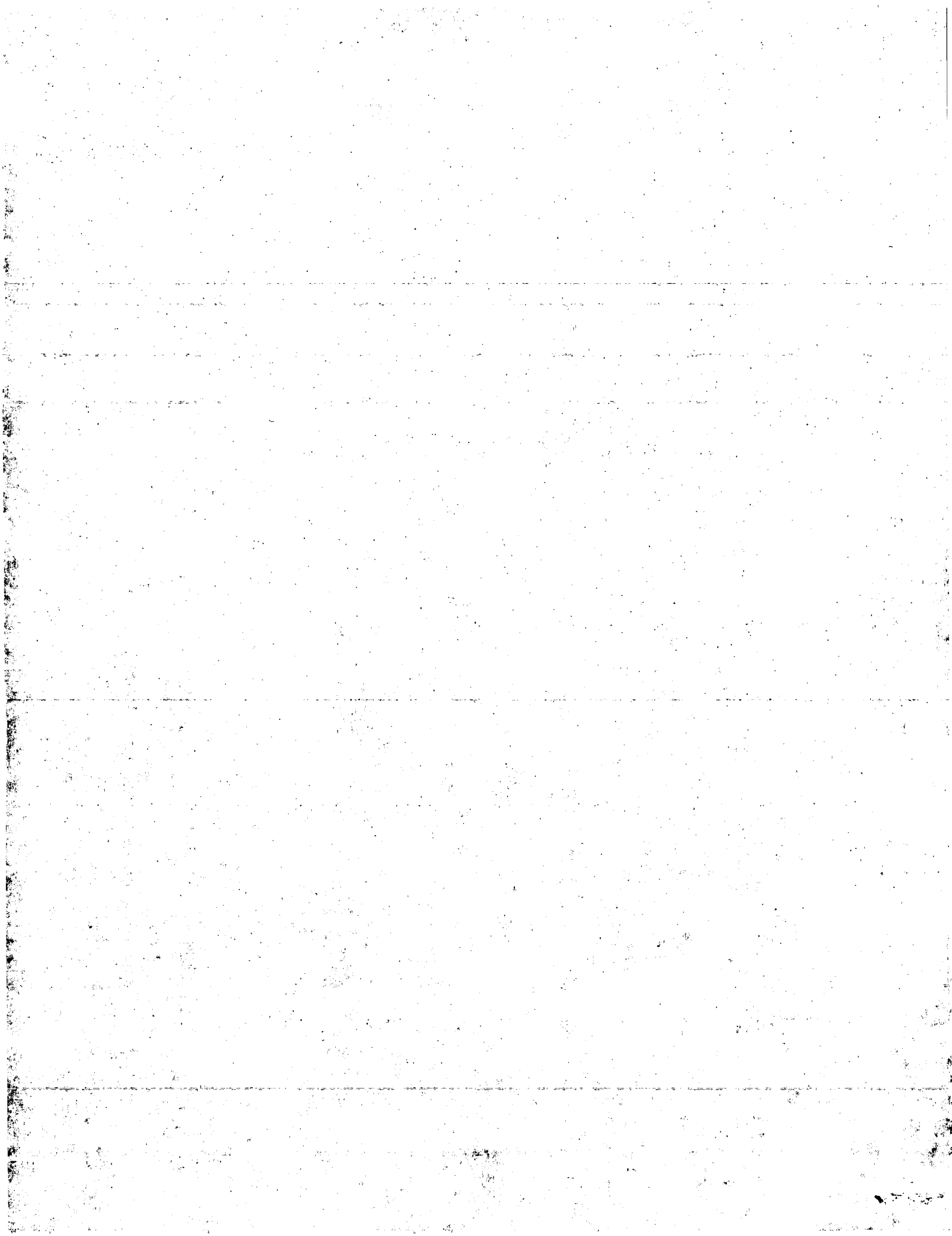
Linear DNA 49 bp PAT 29-MAR-1999

FEATURES

Query Match 1.6%; Score 23.8; DB 6; Length 64;
Best Local Similarity 72.1%; Pred. No. 2.3e+05;

3

Search completed: October 31, 2002, 22:44:37
Job time : 2924 secs



GenCore version 5.1.3
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OM nucleic - nucleic search, using sw model

Run on: October 31, 2002, 21:49:51 ; Search time 59 Seconds
(without alignments)

7568.159 Million cell updates/sec

Title: US-09-919-197-3

Perfect score: 1456

Sequence: 1 tgagcaccagccaccagg.....tatgaacactaaaaaaa 1456

Scoring table:

IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 441362 seqs, 153338381 residues

1 number of hits satisfying chosen parameters: 662272

Minimum DB seq length: 0

Maximum DB seq length: 75

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

Issued_patents_NA.*

1: /cgn2_6/ptodata/1/ina/5A_COMB.seq.*

2: /cgn2_6/ptodata/1/ina/5B_COMB.seq.*

3: /cgn2_6/ptodata/1/ina/6A_COMB.seq.*

4: /cgn2_6/ptodata/1/ina/6B_COMB.seq.*

5: /cgn2_6/ptodata/1/ina/PCTUS_COMB.seq.*

6: /cgn2_6/ptodata/1/ina/backfiles1.seq.*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
C 1	25.4	1.7	72	4	US-09-020-846-51
C 2	23.6	1.6	60	3	US-08-983-607-45
C 3	23.6	1.6	71	1	US-08-487-425-2
C 4	23.6	1.6	71	2	US-08-488-402A-5
C 5	23.6	1.6	71	2	US-08-484-552A-5
C 6	23.6	1.6	71	2	US-08-628-356A-4
C 7	23.6	1.6	71	5	PCT-US96-07439-2
C 8	23.6	1.6	71	5	PCT-US96-09472-5
C 9	22.4	1.5	54	4	US-09-257-581-1
C 10	22.2	1.5	48	4	US-09-425-638A-38
C 11	22.2	1.5	48	4	US-09-543-004-38
C 12	22.2	1.5	61	1	US-07-609-716-22
C 13	22.2	1.5	61	1	US-08-175-155-20
C 14	22.2	1.5	61	1	US-08-477-509B-55
C 15	22.2	1.5	61	1	US-08-707-237A-26
C 16	22.2	1.5	61	3	US-08-482-085B-55
C 17	22.2	1.5	61	3	US-08-475-411A-22
C 18	22.2	1.5	61	4	US-08-478-029A-22
C 19	22.2	1.5	61	4	US-09-444-791A-55
C 20	22.2	1.5	65	1	US-07-609-716-23
C 21	22.2	1.5	65	1	US-08-175-155-21
C 22	22.2	1.5	65	1	US-08-477-509B-56
C 23	22.2	1.5	65	3	US-08-482-085B-56
C 24	22.2	1.5	65	3	US-08-475-411A-23
C 25	22.2	1.5	65	4	US-08-478-029A-23
C 26	22.2	1.5	65	4	US-09-444-791A-56
C 27	22.2	1.5	69	1	US-07-991-867B-29

C 28	22.2	1.5	69	1	US-08-107-755A-29
C 29	22.2	1.5	69	2	US-08-544-332-29
C 30	22.2	1.5	69	4	US-09-370-861A-29
C 31	22	1.5	58	6	5436330-3
C 32	22	1.5	61	1	US-08-055-390-4
C 33	22	1.5	75	4	US-09-042-353-173
C 34	22	1.5	75	4	US-08-758-417A-21
C 35	21.8	1.5	47	4	US-09-641-638-928
C 36	21.8	1.5	71	2	US-08-488-402A-28
C 37	21.8	1.5	71	2	US-08-484-552A-28
C 38	21.8	1.5	71	5	PCT-US96-09472-28
C 39	21.6	1.5	54	2	US-08-412-376-12
C 40	21.6	1.5	54	2	US-08-412-376-26
C 41	21.6	1.5	63	5	PCT-US94-06079-39
C 42	21.6	1.5	70	4	US-09-042-353-172
C 43	21.6	1.5	70	4	US-08-758-417A-20
C 44	21.4	1.5	45	4	US-09-425-638A-13
C 45	21.4	1.5	45	4	US-09-543-004-13

ALIGNMENTS

RESULT 1

US-09-020-846-51/c

; Sequence 51, Application US/09020846

; Patent No. 6322965

; GENERAL INFORMATION:

; APPLICANT: YAMAGUCHI, Kenjiro

; APPLICANT: KASHIWAKUMA, Tomiko

; APPLICANT: CHIBA, Yukie

; APPLICANT: YAGI, Shintaro

; APPLICANT: HASEGAWA, Akira

; TITLE OF INVENTION: CHIMERA ANTIGEN PEPTIDE

; NUMBER OF SEQUENCES: 72

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: FOLEY & LARDNER

; STREET: 3000 K Street, N.W.

; CITY: Washington

; STATE: D.C.

; COUNTRY: U.S.A.

; ZIP: 20007-5109

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; COMPUTER: IBM PC compatible

; OPERATING SYSTEM: PC-DOS/MS-DOS

; SOFTWARE: Patentin Release #1.0, Version #1.30

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/09/020,846

; FILING DATE: 09-FEB-1998

; CLASSIFICATION: 424

; PRIOR APPLICATION DATA:

; APPLICATION NUMBER: JP 9-027015

; FILING DATE: 10-FEB-1997

; PRIOR APPLICATION DATA:

; APPLICATION NUMBER: JP 8-024045

; FILING DATE: 09-FEB-1996

; ATTORNEY/AGENT INFORMATION:

; NAME: Wegner, Harold C.

; REGISTRATION NUMBER: 25,258

; REFERENCE/DOCKET NUMBER: 053466/0225

; TELECOMMUNICATION INFORMATION:

; TELEPHONE: (202) 672-5300

; TELEFAX: (202) 672-5399

; INFORMATION FOR SEQ ID NO: 51:

; SEQUENCE CHARACTERISTICS:

; LENGTH: 72 base pairs

; TYPE: nucleic acid

; STRANDEDNESS: single

; TOPOLOGY: linear

; MOLECULE TYPE: other nucleic acid

; DESCRIPTION: /desc = "Synthetic DNA"

US-09-020-846-51

APPLICANT: JAYASENA, SUMEDHA
APPLICANT: GOLD, LARRY
TITLE OF INVENTION: ENZYME LINKED OLIGONUCLEOTIDE
NUMBER OF SEQUENCES: 3
CORRESPONDENCE ADDRESS:
ADDRESSEE: Swanson & Bratschun, L.L.C. 200
STREET: 8400 E. Prentice Avenue, Suite 200
CITY: Englewood
STATE: Colorado
COUNTRY: USA
ZIP: 80111

Query Match 1.7%; Score 25.4; DB 4; Length 72;
Best Local Similarity 68.6%; Pred. No. 4.4e+02;
Matches 35; Conservative 0; Mismatches 16; Indels 0; Gaps 0;
QY 1081 TGTCTAGTACCTGTTCCAGCCAGGATCAGGATGAGGAGGCTG 1131
Db 70 TGTCTAGTACCTGTTCCAGCTACGTTAAATACCTGGGAGCGGCTG 20

COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3 1/2 diskette, 1.44 MB
COMPUTER: IBM pc compatible
OPERATING SYSTEM: MS-DOS
SOFTWARE: WordPerfect 5.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/487,425
FILING DATE:
CLASSIFICATION: 536
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/714,131
FILING DATE: 10-JUNE-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/536,428
FILING DATE: 11-JUNE-1990
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/964,624
FILING DATE: 21-OCTOBER-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/234,997
FILING DATE: 28-APRIL-1994
ATTORNEY/AGENT INFORMATION:
NAME: Barry J. Swanson
REGISTRATION NUMBER: 33,215
REFERENCE/DOCKET NUMBER:
TELEPHONE: (303) 793-3333
TELEFAX: (303) 793-3433
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 71 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: RNA
FEATURE:
OTHER INFORMATION: All C's are 2'-NH2 cytosine
FEATURE:
OTHER INFORMATION: All U's are 2'-NH2 uracil
US-08-487-425-2

Query Match 1.7%; Score 25.4; DB 4; Length 72;
Best Local Similarity 68.6%; Pred. No. 4.4e+02;
Matches 35; Conservative 0; Mismatches 16; Indels 0; Gaps 0;
QY 1081 TGTCTAGTACCTGTTCCAGCCAGGATCAGGATGAGGAGGCTG 1131
Db 70 TGTCTAGTACCTGTTCCAGCTACGTTAAATACCTGGGAGCGGCTG 20

COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5" 1.44 MB diskette
COMPUTER: IBM PC
OPERATING SYSTEM: MS DOS
SOFTWARE: Word Processing
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/983,607
FILING DATE: April 27, 1998
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/IB96/01032
FILING DATE: June 28, 1996
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Mary M. Krinsky
REGISTRATION NUMBER: 32423
REFERENCE/DOCKET NUMBER: OCR-679
TELEPHONE: 203-773-9544
TELEFAX: 203-773-1183
INFORMATION FOR SEQ ID NO: 45:
SEQUENCE CHARACTERISTICS:
LENGTH: 60 residues
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
DESCRIPTION: primer used in constructs
US-08-983-607-45

Query Match 1.6%; Score 23.6; DB 3; Length 60;
Best Local Similarity 66.7%; Pred. No. 1.3e+03;
Matches 32; Conservative 1; Mismatches 15; Indels 0; Gaps 0;
QY 694 TTACCTCTCTAAAGGAGGAGTGTAGTGGGAGCTCAAGGCGGAGC 741
Db 50 TCACCTCTCTTAAGGAGGAGTGTAGTGGGAGCTCAAGGCGGAGC 3

COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3 1/2 diskette, 1.44 MB
COMPUTER: IBM pc compatible
OPERATING SYSTEM: MS-DOS
SOFTWARE: WordPerfect 5.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/487,425
FILING DATE:
CLASSIFICATION: 536
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/714,131
FILING DATE: 10-JUNE-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/536,428
FILING DATE: 11-JUNE-1990
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/964,624
FILING DATE: 21-OCTOBER-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/234,997
FILING DATE: 28-APRIL-1994
ATTORNEY/AGENT INFORMATION:
NAME: Barry J. Swanson
REGISTRATION NUMBER: 33,215
REFERENCE/DOCKET NUMBER:
TELEPHONE: (303) 793-3333
TELEFAX: (303) 793-3433
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 71 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: RNA
FEATURE:
OTHER INFORMATION: All C's are 2'-NH2 cytosine
FEATURE:
OTHER INFORMATION: All U's are 2'-NH2 uracil
US-08-487-425-2

Query Match 1.6%; Score 23.6; DB 3; Length 60;
Best Local Similarity 66.7%; Pred. No. 1.3e+03;
Matches 32; Conservative 1; Mismatches 15; Indels 0; Gaps 0;
QY 694 TTACCTCTCTAAAGGAGGAGTGTAGTGGGAGCTCAAGGCGGAGC 741
Db 50 TCACCTCTCTTAAGGAGGAGTGTAGTGGGAGCTCAAGGCGGAGC 3

RESULT 4
US-08-488-402A-5/C
Sequence 5, Application US/08488402A
Patent No. 5837456
GENERAL INFORMATION:
APPLICANT: GOLD ET AL.
TITLE OF INVENTION: HIGH AFFINITY OLIGONUCLEOTIDE LIGANDS TO
TITLE OF INVENTION: CHORIONIC GONADOTROPIN HORMONE AND RELATED GLYCOPROTEIN
TITLE OF INVENTION: HORMONES
NUMBER OF SEQUENCES: 160
CORRESPONDENCE ADDRESS:

RESULT 3
US-08-487-425-2/C
Sequence 2, Application US/08487425
Patent No. 5789163
GENERAL INFORMATION:
APPLICANT: DROLET, DAN

;; FILING DATE: 07 JUNE 1995
;; PRIORITY APPLICATION DATA:
;; APPLICATION NUMBER: 08/484,552
;; FILING DATE: 07 JUNE 1995
;; ATTORNEY/AGENT INFORMATION:
;; NAME: Barry J. Swanson
;; REGISTRATION NUMBER: 33,215
;; REFERENCE/DOCKET NUMBER: NEX36-1
;; TELECOMMUNICATION INFORMATION:
;; TELEPHONE: (303) 793-3333
;; TELEFAX: (303) 793-3433
;; INFORMATION FOR SEQ ID NO: 5:
;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 71 base pairs
;; TYPE: nucleic acid
;; STRANDEDNESS: single
;; TOPOLOGY: linear
;; FEATURE:
;; OTHER INFORMATION: All C's are 2'-NH2 modified
;; FEATURE:
;; OTHER INFORMATION: All U's are 2'-NH2 modified
US96-09472-5

Query Match 1.6%; Score 23.6; DB 5; Length 71;
Best Local Similarity 69.6%; Pred. No. 1.4e+03;
Matches 32; Conservative 0; Mismatches 14; Indels 0; Gaps 0;

QY 952 GCGCGCTGACCGCTGCTCTCCACGGCTCCACCTCAAGTCCAT 997
DB 63 GTCGCTGCCAGTCCCTCCACGTGCCCTCCACACTCAGGCCCTT 18

RESULT 9

US-09-257-581-1/c
; Sequence 1, Application US/09257581
; Patent No. 6207419

;; GENERAL INFORMATION:
;; APPLICANT: Church, Frank C.
;; APPLICANT: Bauman, Susannah J.
;; TITLE OF INVENTION: THROMBIN INHIBITORY AGENTS AND METHODS OF USING SAME
;; FILE REFERENCE: 5470-232
;; CURRENT APPLICATION NUMBER: US/09/257,581
;; CURRENT FILING DATE: 1999-02-25
;; EARLIER APPLICATION NUMBER: 60/076,210
;; EARLIER FILING DATE: 1998-02-27
;; NUMBER OF SEQ ID NOS: 7
;; SOFTWARE: PatentIn Ver. 2.0
;; SEQ ID NO 1
;; LENGTH: 54
;; TYPE: DNA
;; ORGANISM: Artificial Sequence
;; FEATURE:
;; OTHER INFORMATION: Description of Artificial Sequence: primer
US-09-257-581-1

Query Match 1.5%; Score 22.4; DB 4; Length 54;
Best Local Similarity 72.5%; Pred. No. 2.6e+03;
Matches 29; Conservative 0; Mismatches 11; Indels 0; Gaps 0;

QY 690 TAAATTAATCTCTCTAAAGGAGGAGGTAGTAGTGGACCT 729
DB 52 TAGACCTCCACCTCTAGTGGTGGTGGTGGGACCT 13

RESULT 10

US-09-425-638A-38
; Sequence 38, Application US/09425638A
; Patent No. 6342587

;; GENERAL INFORMATION:
;; APPLICANT: Carlos F. Barbas III, Christoph Rader, Gerd Ritter, Sydney Welt and
;; APPLICANT: Lloyd J. Old
;; TITLE OF INVENTION: A33 ANTIGEN SPECIFIC IMMUNOGLOBULIN PRODUCTS AND USES THEREOF
;; FILE REFERENCE: LUD 5630

;; CURRENT APPLICATION NUMBER: US/09/425.638A
;; CURRENT FILING DATE: 1999-10-22
;; NUMBER OF SEQ ID NOS: 129
;; SEQ ID NO 38
;; LENGTH: 48
;; TYPE: DNA
;; ORGANISM: Homo sapiens
;; FEATURE:
US-09-425-638A-38

Query Match 1.5%; Score 22.2; DB 4; Length 48;
Best Local Similarity 69.8%; Pred. No. 2.8e+03;
Matches 30; Conservative 0; Mismatches 13; Indels 0; Gaps 0;

QY 309 GCTGTGACCTTTGAGTGGCTGAGGCCCGCGGTGCCACGATAC 351
DB 5 GATGGGCCCTTGGTGAGGCTGAGGAGAGCGGTGACCGGTGC 47

RESULT 11

US-09-543-004-38
; Sequence 38, Application US/09543004
; Patent No. 6346249

;; GENERAL INFORMATION:
;; APPLICANT: Carlos F. Barbas III, Christoph Rader, Gerd Ritter, Sydney Welt and
;; APPLICANT: Lloyd J. Old
;; TITLE OF INVENTION: A33 ANTIGEN SPECIFIC IMMUNOGLOBULIN PRODUCTS AND USES THE
;; FILE REFERENCE: LUD 5630.1
;; CURRENT APPLICATION NUMBER: US/09/543,004
;; CURRENT FILING DATE: 2000-04-04
;; PRIOR APPLICATION NUMBER: 09/425,638
;; PRIOR FILING DATE: 1999-10-22
;; NUMBER OF SEQ ID NOS: 129
;; SEQ ID NO 38
;; LENGTH: 48
;; TYPE: DNA
;; ORGANISM: Homo sapiens
;; FEATURE:
US-09-543-004-38

Query Match 1.5%; Score 22.2; DB 4; Length 48;
Best Local Similarity 69.8%; Pred. No. 2.8e+03;
Matches 30; Conservative 0; Mismatches 13; Indels 0; Gaps 0;

QY 309 GCTGTGACCTTTGAGTGGCTGAGGCCCGCGGTGCCACGATAC 351
DB 5 GATGGGCCCTTGGTGAGGCTGAGGAGAGCGGTGACCGGTGC 47

RESULT 12

US-07-609-716-22/c
; Sequence 22, Application US/07609716
; Patent No. 5514581

;; GENERAL INFORMATION:
;; APPLICANT: Ferrari, Franco A.
;; APPLICANT: Cappello, Joseph
;; TITLE OF INVENTION: Functional Recombinantly Prepared
;; TITLE OF INVENTION: Synthetic Protein Polymer
;; NUMBER OF SEQUENCES: 118
;; CORRESPONDENCE ADDRESS:
;; ADDRESSEE: Flehr, Hohbach, Test, Albritton & Herbert
;; STREET: Four Embarcadero Center, Suite 3400
;; CITY: San Francisco
;; STATE: CA
;; COUNTRY: US
;; ZIP: 94111
;; COMPUTER READABLE FORM:
;; MEDIUM TYPE: Floppy disk
;; COMPUTER: IBM PC compatible
;; OPERATING SYSTEM: PC-DOS/MS-DOS
;; SOFTWARE: PatentIn Release #1.0, Version #1.30
;; CURRENT APPLICATION DATA:
;; APPLICATION NUMBER: US/07/609,716

RESULT 13
 US-08-175-155-20/C
 : Sequence 20, Application US/08175155
 : Patent No. 5641648
 : GENERAL INFORMATION:
 : APPLICANT: Ferrazi, Franco A.
 : APPLICANT: Cappello, Joseph
 : APPLICANT: Crissman, John W.
 : APPLICANT: Dorman, Mary A.
 : TITLE OF INVENTION: Methods for Preparing Synthetic
 : TITLE OF INVENTION: Repetitive DNA
 : NUMBER OF SEQUENCES: 69
 : CORRESPONDENCE ADDRESS:
 : ADDRESSEE: Flehr, Hohbach, Test, Albritton & Herbert
 : STREET: Four Embarcadero Center, Suite 3400
 : CITY: San Francisco
 : STATE: CA
 : COUNTRY: US
 : ZIP: 94111
 : COMPUTER READABLE FORM:
 : MEDIUM TYPE: Floppy disk
 : COMPUTER: IBM PC compatible
 : OPERATING SYSTEM: PC-DOS/MS-DOS
 : SOFTWARE: PatentIn Release #1.0, Version #1.30
 : CURRENT APPLICATION DATA:
 : APPLICATION NUMBER: US/08/175,155
 : FILING DATE: 29-DEC-1993
 : CLASSIFICATION: 435
 : ATTORNEY/AGENT INFORMATION:
 : NAME: Rowland, Bertram I.
 : REGISTRATION NUMBER: 20015
 : REFERENCE/DOCKET NUMBER: A-55186-5/BIR
 : TELECOMMUNICATION INFORMATION:
 : TELEPHONE: 415-781-1989
 : TELEFAX: 415-398-3249
 : INFORMATION FOR SEQ ID NO: 20:
 : SEQUENCE CHARACTERISTICS:
 : LENGTH: 61 base pairs
 : TYPE: nucleic acid
 : STRANDEDNESS: single
 : TOPOLOGY: linear
 : MOLECULE TYPE: cdna
 : US-08-175-155-20

Query Match 1.5%; Score 22.2; DB 1; Length 61;
 Best Local Similarity 64.7%; Pred. No. 3.2e+03;
 Matches 33; Conservative 0; Mismatches 18; Indels 0; Gaps 0;

Qy	229	GGCAGCTGCTCCCGAGGACGCGGCTGCTGCAGGGTCTCGGGGCC	279
Db	51	GCCAGAGCCCGCGCCAGCTCCAGAACGGCTCTCGACCGCTCCCGGCACC	1

US-08-175-155-20

RESULT 15

	Query Match	1.5%	Score	22.2;	DB 2;	Length	61;
	Best Local Similarity	64.7%;	Pred. No.	3.2e+03;			
	Matches	33;	Conservative	0;	Mismatches	18;	Indels
							Gaps
							0;
QY	229	GGCAGCTGCTCC	CCAGGAC	CACGCGGCTGCTG	CAGGTTT	TGCTGGGGCC	279
		1 1 1	1 1 1	1 1 1	1 1 1	1 1 1	1
DB	51	GCAGAGCCCG	CGCCAGCTC	CAGAACC	GGCTCTG	CACCGCTCGCGG	1
		1 1 1	1 1 1	1 1 1	1 1 1	1 1 1	1

Search completed: October 31, 2002, 23:18:35
Job time : 61 secs

GenCore version 5.1.3
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OM nucleic - nucleic search, using sw model

Run on: October 31, 2002, 21:51:41 ; Search time 57 Seconds
(without alignments)
8510.060 Million cell updates/sec

Title: US-09-919-197-3
Perfect score: 1456
Sequence: 1 tgagcaccagccacacagg.....tatgaacactataaaaaa 1456

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 310279 seqs, 166577418 residues
1 number of hits satisfying chosen parameters: 169348

Minimum DB seq length: 0
Maximum DB seq length: 75

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : Published_Applications_NA:
1: /cgn2_6/ptodata/1/pubpna/PCTUS_PUBCOMB.seq.*
2: /cgn2_6/ptodata/1/pubpna/PCT_NEW_PUB.seq.*
3: /cgn2_6/ptodata/1/pubpna/US06_NEW_PUB.seq.*
4: /cgn2_6/ptodata/1/pubpna/US06_PUBCOMB.seq.*
5: /cgn2_6/ptodata/1/pubpna/US07_NEW_PUB.seq.*
6: /cgn2_6/ptodata/1/pubpna/PCTUS_PUBCOMB.seq.*
7: /cgn2_6/ptodata/1/pubpna/US08_NEW_PUB.seq.*
8: /cgn2_6/ptodata/1/pubpna/US08_PUBCOMB.seq.*
9: /cgn2_6/ptodata/1/pubpna/US09_NEW_PUB.seq.*
10: /cgn2_6/ptodata/1/pubpna/US09_PUBCOMB.seq.*
11: /cgn2_6/ptodata/1/pubpna/US10_NEW_PUB.seq.*
12: /cgn2_6/ptodata/1/pubpna/US10_PUBCOMB.seq.*
13: /cgn2_6/ptodata/1/pubpna/US60_NEW_PUB.seq.*
14: /cgn2_6/ptodata/1/pubpna/US60_PUBCOMB.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	26	1.8	73	10	US-09-466-320-16
2	21	1.4	56	10	US-09-944-036-30
3	21	1.4	64	10	US-09-919-580-637
4	21	1.4	74	10	US-09-250-611-7
5	20.6	1.4	70	10	US-09-878-574-8527
6	20.4	1.4	55	10	US-09-790-417-19
7	20.4	1.4	67	10	US-09-878-574-5162
8	20.4	1.4	71	10	US-09-783-590-2509
9	20.4	1.4	73	10	US-09-783-590-3398
10	20.2	1.4	69	10	US-09-983-965-3899
11	20	1.4	51	10	US-09-874-547-8
12	20	1.4	70	10	US-09-783-590-4790
13	20	1.4	75	10	US-09-864-761-28813
14	19.8	1.4	60	10	US-09-925-301-758
15	19.8	1.4	60	10	US-09-975-408-68
16	19.8	1.4	60	12	US-10-075-579-68
17	19.8	1.4	70	10	US-09-822-250-30
18	19.8	1.4	72	10	US-09-884-441-304
19	19.8	1.4	73	12	US-10-108-280-18

c	20	19.8	1.4	75	10	US-09-922-261-151	Sequence 151, Appl
	21	19.6	1.3	48	10	US-09-810-502-29	Sequence 29, Appl
	22	19.6	1.3	55	10	US-09-874-547-48	Sequence 48, Appl
	23	19.6	1.3	56	9	US-09-144-886-39	Sequence 39, Appl
	24	19.6	1.3	56	10	US-09-874-547-44	Sequence 44, Appl
	25	19.6	1.3	56	10	US-09-988-899-49	Sequence 49, Appl
c	26	19.6	1.3	60	10	US-09-815-242-1130	Sequence 1130, Ap
c	27	19.6	1.3	71	10	US-09-783-590-5311	Sequence 5311, Ap
	28	19.6	1.3	73	10	US-09-896-096A-4	Sequence 4, Appli
	29	19.6	1.3	73	10	US-09-894-924-4	Sequence 4, Appli
c	30	19.4	1.3	55	10	US-09-920-300A-1151	Sequence 1151, Ap
c	31	19.4	1.3	55	12	US-10-033-528-1151	Sequence 1151, Ap
	32	19.4	1.3	67	10	US-09-943-286-6	Sequence 6, Appli
c	33	19.2	1.3	26	10	US-09-780-172-13	Sequence 13, Appl
c	34	19.2	1.3	45	10	US-09-989-722-122	Sequence 122, App
c	35	19.2	1.3	45	10	US-09-989-723-122	Sequence 122, App
c	36	19.2	1.3	45	10	US-09-989-279-122	Sequence 122, App
c	37	19.2	1.3	45	10	US-09-989-727-122	Sequence 122, App
c	38	19.2	1.3	45	10	US-09-989-731-122	Sequence 122, App
c	39	19.2	1.3	45	10	US-09-989-732-122	Sequence 122, App
c	40	19.2	1.3	45	10	US-09-991-073-122	Sequence 122, App
c	41	19.2	1.3	45	10	US-09-990-442-122	Sequence 122, App
c	42	19.2	1.3	45	10	US-09-991-163-122	Sequence 122, App
c	43	19.2	1.3	45	10	US-09-993-604-122	Sequence 122, App
c	44	19.2	1.3	45	10	US-09-990-456-122	Sequence 122, App
c	45	19.2	1.3	45	10	US-09-989-721-122	Sequence 122, App

ALIGNMENTS

RESULT 1
US-09-466-320-16
; Sequence 16, Application US/09466320
; Patent No. US20020025939A1
; GENERAL INFORMATION:
; APPLICANT: Iversen, Patrick
; TITLE OF INVENTION: Chorionic Gonadotropin DNA Vaccines and
; FILE REFERENCE: 0450-0026.30
; CURRENT APPLICATION NUMBER: US/09/466,320
; CURRENT FILING DATE: 1999-12-17
; EARLIER APPLICATION NUMBER: US 60/112,910
; EARLIER FILING DATE: 1998-12-18
; NUMBER OF SEQ ID NOS: 25
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 16
; LENGTH: 73
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: coding sequence
US-09-466-320-16

Query Match	1.8%	Score 26;	DB 10;	Length 73;
Best Local Similarity	62.1%	Pred. No. 3.4e+02;		
Matches	41;	Conservative	0;	Mismatches 25;
				Indels 0;
				Gaps 0;
QY	186	CTGGCCACAGAGTGGCTTCTCTAGAACCTGCGATCCTCTGGCAGCTGCTCCCCAG	245	
Db	8	CCGTGCTAGCAATGAGACCGCTGAACAATACCAACCCCTGTACAGGGGCTCCCCAG	67	
QY	246	GACCAAG 251		
Db	68	GATAAG 73		
RESULT 2				
US-09-944-036-30				
; Sequence 30, Application US/09944036				
; Patent No. US2002005095A1				
; GENERAL INFORMATION:				
; APPLICANT: YANG, Yeasing Y.				

APPLICANT: BRENTANO, Steven T.
APPLICANT: BABOLA, Odile
APPLICANT: TRAN, Nathalie
APPLICANT: VERNET, Guy
TITLE OF INVENTION: AMPLIFICATION OF HIV-1 SEQUENCES FOR DETECTION OF
SEQUENCES ASSOCIATED WITH DRUG-RESISTANCE MUTATIONS
FILE REFERENCE: GP114-02.UT
CURRENT APPLICATION NUMBER: US/09/944,036
CURRENT FILING DATE: 2001-08-31
PRIOR APPLICATION NUMBER: US 60/229,790
PRIOR FILING DATE: 2000-09-01
NUMBER OF SEQ ID NOS: 70
SOFTWARE: PatentIn Ver. 2.1
SEQ ID NO 30
LENGTH: 56
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Description of Artificial Sequence: Capture
9-944-036-30
Query Match 1.4%; Score 21; DB 10; Length 56;
Best Local Similarity 62.3%; Pred. No. 7.6e+03;
Matches 33; Conservative 0; Mismatches 20; Indels 0; Gaps 0;
QY 1404 CTGAATTAAGTATTGATTTTGTAAATAAAGGTATGAACACTAAAAA 1456
Db 2 CTGGAATAACTTCGCTCTATTATTTAAAAA 1456
RESULT 3
US-09-919-580-637/c
Sequence 637, Application US/09919580
Patent No. US20020110832A1
GENERAL INFORMATION:
APPLICANT: Xu, Jiangchun
APPLICANT: Secret, Heather
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY
TITLE OF INVENTION: AND DIAGNOSIS OF COLON CANCER
FILE REFERENCE: 210121.552
CURRENT APPLICATION NUMBER: US/09/919,580
CURRENT FILING DATE: 2001-07-30
NUMBER OF SEQ ID NOS: 934
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 637
LENGTH: 64
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: misc_feature
LOCATION: 56
OTHER INFORMATION: n = A,T,C or G
US-09-919-580-637
Query Match 1.4%; Score 21; DB 10; Length 64;
Best Local Similarity 65.2%; Pred. No. 8.2e+03;
Matches 30; Conservative 0; Mismatches 16; Indels 0; Gaps 0;
QY 1411 AAGTTATGATTTTGTATAAAGGTATGAACACTAAAAA 1456
Db 64 AAGCGTTTCTTTTAAAAA 1456
RESULT 4
US-09-250-611-7/c
Sequence 7, Application US/09250611
Patent No. US20020143161A1
GENERAL INFORMATION:
APPLICANT: Byrne, Jennifer A.
APPLICANT: Basset, Paul
TITLE OF INVENTION: Members of the D52 Gene Family

FILE REFERENCE: 1363.0210001
CURRENT APPLICATION NUMBER: US/09/250,611
CURRENT FILING DATE: 1999-02-17
NUMBER OF SEQ ID NOS: 108
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 7
LENGTH: 74
TYPE: DNA
ORGANISM: Homo sapiens
US-09-250-611-7
Query Match 1.4%; Score 21; DB 10; Length 74;
Best Local Similarity 62.3%; Pred. No. 9e+03;
Matches 33; Conservative 0; Mismatches 20; Indels 0; Gaps 0;
QY 323 GGTGGGTGAGGCCCGCTGCCAGCATACATCAAGAAGATTCTGCTGGAGGAGC 375
Db 74 GGTCCGAGAGTTCTCTCATGCTGTCATATTGAGTGGCGATGGAGTAGC 22
RESULT 5
US-09-878-574-8527/c
Sequence 8527, Application US/09878574
Patent No. US20020110548A1
GENERAL INFORMATION:
APPLICANT: Byrum, Joseph R.
APPLICANT: La Rosa, Thomas J.
APPLICANT: Thompson, Michael D.
TITLE OF INVENTION: Nucleic Acid Molecules and Other Molecules Associated with
TITLE OF INVENTION: Plants
FILE REFERENCE: 38-21(15401)B
CURRENT APPLICATION NUMBER: US/09/878,574
CURRENT FILING DATE: 2001-12-21
PRIOR APPLICATION NUMBER: 09/333,535
PRIOR FILING DATE: 1999-06-14
NUMBER OF SEQ ID NOS: 15775
SEQ ID NO 8527
LENGTH: 70
TYPE: DNA
ORGANISM: Glycine max
OTHER INFORMATION: Clone ID: 701101333H1
US-09-878-574-8527
Query Match 1.4%; Score 20.6; DB 10; Length 70;
Best Local Similarity 59.3%; Pred. No. 1.1e+04;
Matches 35; Conservative 0; Mismatches 24; Indels 0; Gaps 0;
QY 1355 AAGCTTAGGCTGCTTATTGGACAGAGTCTCTATGACTTTATACAGAACTGAATTAAG 1413
Db 68 AGGCTGGCGCTCATCATTCGACCGCGCTTCCCAAGGAGACCTTCCCGGAATCGGTTTCGAG 10
RESULT 6
US-09-790-417-19
Sequence 19, Application US/09790417
Patent No. US20010031470A1
GENERAL INFORMATION:
APPLICANT: Shultz, John W.
APPLICANT: Lewis, Martin K.
APPLICANT: Lieppe, Donna
APPLICANT: Mandrekar, Michelle
APPLICANT: Kephart, Daniel
APPLICANT: Rhodes, Richard B.
APPLICANT: Andrews, Christine A.
APPLICANT: Hartnett, James R.
APPLICANT: Gu, Trent
APPLICANT: Olson, Ryan J.
APPLICANT: Wood, Keith W.
APPLICANT: Welch, Roy
APPLICANT: Nucleic Acid Detection
TITLE OF INVENTION: PRO-103 6868/75528
FILE REFERENCE: PRO-103 6868/75528
CURRENT APPLICATION NUMBER: US/09/790,417
CURRENT FILING DATE: 2001-02-22


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; PRIOR APPLICATION NUMBER: 09/358,972
; PRIOR FILING DATE: 1999-07-21
; PRIOR APPLICATION NUMBER: 09/042,287
; PRIOR FILING DATE: 1998-03-13
; NUMBER OF SEQ ID NOS: 290
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 19
; LENGTH: 55
; TYPE: DNA
; ORGANISM: Cytomegalovirus
; FEATURE:
; OTHER INFORMATION: target for mutant cytomegalovirus
US-09-790-417-19

Query Match          1.4%; Score 20.4; DB 10; Length 55;
Best Local Similarity 61.1%; Pred. No. 1.1e+04;
Matches 33; Conservative 0; Mismatches 21; Indels 0; Gaps 0;

QY 8 CAGCCAACGAGGGCTGCCATGCCAGGAGCTGCAAGCGCCGCCCATCTT 61
      ||||| | ||||| | ||||| | ||||| | ||||| | ||||| |
1 CAGCAGTGGCTGAGCTTCCGGTCTCCGGCGGCGGCGGCGCGCGGTGCT 54

RESULT 7
US-09-878-574-5162
; Sequence 5162, Application US/09878574
; Patent No. US20020110548A1
; GENERAL INFORMATION:
; APPLICANT: Byrum, Joseph R.
; APPLICANT: La Rosa, Thomas J.
; APPLICANT: Thompson, Michael D.
; TITLE OF INVENTION: Nucleic Acid Molecules and Other Molecules Associated with
; FILE OF INVENTION: Plants
; FILE REFERENCE: 38-21(15401)B
; CURRENT APPLICATION NUMBER: US/09/878,574
; CURRENT FILING DATE: 2001-12-21
; PRIOR APPLICATION NUMBER: 09/333,535
; PRIOR FILING DATE: 1999-06-14
; NUMBER OF SEQ ID NOS: 15775
; SEQ ID NO 5162
; LENGTH: 67
; TYPE: DNA
; ORGANISM: Glycine max
; OTHER INFORMATION: Clone ID: LIB3028-022-Q1-B1-D9
US-09-878-574-5162

Query Match          1.4%; Score 20.4; DB 10; Length 67;
Best Local Similarity 61.1%; Pred. No. 1.3e+04;
Matches 33; Conservative 0; Mismatches 21; Indels 0; Gaps 0;

QY 1403 ACTGAATTAAGTTATTTGTTTGAATAAAGGTATGAACACTTAAAAA 1456
      || ||||| | ||||| | ||||| | ||||| | ||||| | ||||| |
DB 11 ACCCAATAGTGAACCTTTTGTGCAAAAAAATAAATAAATAAACA 64

RESULT 8
US-09-783-590-3398
; Sequence 3398, Application US/09783590
; Patent No. US20020110850A1
; GENERAL INFORMATION:
; APPLICANT: Dillon, Patrick J.
; APPLICANT: Haseltine, William A.
; APPLICANT: Li, Haodong
; APPLICANT: Rosen, Craig A.
; APPLICANT: Ruben, Steven M.
; TITLE OF INVENTION: Human Genes, Sequences, and Expression Products 16.2
; FILE REFERENCE: PO-16.2C1
; CURRENT APPLICATION NUMBER: US/09/783,590
; CURRENT FILING DATE: 2000-02-15
; PRIOR APPLICATION NUMBER: 08/420,856
; PRIOR FILING DATE: 1995-04-12
; PRIOR FILING DATE: 1994-11-21
; NUMBER OF SEQ ID NOS: 12485
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 3398
; LENGTH: 73
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (32)
; OTHER INFORMATION: n equals a,t,g, or c
US-09-783-590-3398

Query Match          1.4%; Score 20.4; DB 10; Length 73;
Best Local Similarity 54.9%; Pred. No. 1.3e+04;
Matches 39; Conservative 0; Mismatches 32; Indels 0; Gaps 0;
```

```
; NUMBER OF SEQ ID NOS: 12485
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 2509
; LENGTH: 71
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (19)
; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc feature
; LOCATION: (46)
; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc feature
; LOCATION: (54)
; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc feature
; LOCATION: (63)
; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc feature
; LOCATION: (64)
; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc feature
; LOCATION: (67)
; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc feature
; LOCATION: (68)
; OTHER INFORMATION: n equals a,t,g, or c
US-09-783-590-2509

Query Match          1.4%; Score 20.4; DB 10; Length 71;
Best Local Similarity 56.6%; Pred. No. 1.3e+04;
Matches 30; Conservative 0; Mismatches 23; Indels 0; Gaps 0;

QY 1404 CTGAATTAAGTTATTTGTTTGAATAAAGGTATGAACACTTAAAAA 1456
      | ||| | | ||||| ||||| | ||||| | ||||| | ||||| |
DB 69 CNAANNAAAAAAANTTTTTTNAAAAAAATAAATAAATAAATAA 17

RESULT 9
US-09-783-590-3398
; Sequence 3398, Application US/09783590
; Patent No. US20020110850A1
; GENERAL INFORMATION:
; APPLICANT: Dillon, Patrick J.
; APPLICANT: Haseltine, William A.
; APPLICANT: Li, Haodong
; APPLICANT: Rosen, Craig A.
; APPLICANT: Ruben, Steven M.
; TITLE OF INVENTION: Human Genes, Sequences, and Expression Products 16.2
; FILE REFERENCE: PO-16.2C1
; CURRENT APPLICATION NUMBER: US/09/783,590
; CURRENT FILING DATE: 2000-02-15
; PRIOR APPLICATION NUMBER: 08/420,856
; PRIOR FILING DATE: 1995-04-12
; PRIOR FILING DATE: 1994-11-21
; NUMBER OF SEQ ID NOS: 12485
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 3398
; LENGTH: 73
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (32)
; OTHER INFORMATION: n equals a,t,g, or c
US-09-783-590-3398

Query Match          1.4%; Score 20.4; DB 10; Length 73;
Best Local Similarity 54.9%; Pred. No. 1.3e+04;
Matches 39; Conservative 0; Mismatches 32; Indels 0; Gaps 0;
```

Query Match	1.48;	Score 20;	DB 10;	Length 51;
Best Local Similarity	72.28;	Pred. No. 1.4e+04;		

```

RESULT 13
US - 98-864-761-28813/c
; Sequence 28813, Application US/09864761
; Patent No. US20020048763A1
; GENERAL INFORMATION:
; APPLICANT: Penn, Sharon G.
; APPLICANT: Rank, David R.
; APPLICANT: Hanzel, David K.
; APPLICANT: Chen, Wensheng
; TITLE OF INVENTION: HUMAN GENOME-DERIV
; TITLE OF INVENTION: GENE EXPRESSION A
; FILE REFERENCE: Acomica-x-1

```


GenCore version 5.1.3
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OM nucleic - nucleic search, using sw model

Run on: October 31, 2002, 21:09:57 ; Search time 1937 Seconds
(without alignments)
12173.789 Million cell updates/sec

Title: US-09-919-197-3

Perfect score: 1456

Sequence: 1 tgaacaccagccaccagg.....tatgaacactaaaaaaa 1456

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 16154066 seqs, 8097743376 residues

al number of hits satisfying chosen parameters: 218680

Minimum DB seq length: 0

Maximum DB seq length: 75

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

EST:*

1: em_estba:**
2: em_esthum:**
3: em_estin:**
4: em_estmu:**
5: em_estov:**
6: em_estpl:**
7: em_estro:**
8: em_htc:**
9: gb_est1:**
10: gb_est2:**
11: gb_htc:**
12: gb_est3:**
13: gb_est4:**
14: gb_est5:**
15: em_estfun:**
16: em_estom:**
17: gb_gss:**
18: em_gss_hum:**
19: em_gss_inv:**
20: em_gss_pln:**
21: em_gss_vrt:**
22: em_gss_fun:**
23: em_gss_mam:**
24: em_gss_mus:**
25: em_gss_othr:**
26: em_gss_pro:**
27: em_gss_rod:**

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
C 1	32.4	2.2	34	9	A1221747
C 2	29.8	2.0	62	17	B37921
C 3	28.8	2.0	57	13	BI745330
C 4	28.2	1.9	52	9	AA782923
C 5	27.6	1.9	72	13	BI260363
C 6	26.8	1.8	68	12	BF632727

C	7	26.6	1.8	53	14	H55352	H55352 CHR220291 C
	8	26.2	1.8	35	14	R37139	R37139 yf54g11.s1
	9	26.2	1.8	55	9	AI123105	AI123105 qa85b11.s
	10	26.2	1.8	60	14	R85541	R85541 y038c03.s1
	11	25.8	1.8	47	14	H97244	H97244 yv98e05.s1
	12	25.6	1.8	46	9	AA916309	AA916309 on64a06.s
	13	25.4	1.7	61	12	BF581452	BF581452 602100856
C	14	25.4	1.7	63	10	AW333106	AW333106 S17D5 AGS
	15	25.2	1.7	47	14	N67514	N67514 za08g10.s1
	16	25.2	1.7	69	17	CNS0214K	AL198461 Tetraodon
	17	25	1.7	65	17	AZ664880	AZ664880 LM0545H01
	18	25	1.7	74	9	AA574536	AA574536 vm29e03.r
C	19	24.8	1.7	46	9	AA853962	AA853962 aj51e02.s
C	20	24.8	1.7	61	9	AI144267	AI144267 qb59e02.x
C	21	24.8	1.7	67	10	AW633089	AW633089 bl03f09.x
C	22	24.8	1.7	67	17	AZ397521	AZ397521 LM0162J05
C	23	24.8	1.7	75	14	N68192	N68192 zal1f07.s1
C	24	24.6	1.7	67	9	AI366713	AI366713 qy95b04.x
C	25	24.6	1.7	73	9	AA987684	AA987684 q088a08.s
C	26	24.4	1.7	70	14	BQ454399	BQ454399 ke04b11.y
C	27	24.2	1.7	63	9	AI274192	AI274192 ql46b08.x
C	28	24.2	1.7	71	14	T25646	T25646 EST00515 Eq
C	29	24	1.6	64	9	AI448858	AI448858 ms45f07.x
C	30	24	1.6	69	17	CNS0214K	AL198461 Tetraodon
C	31	24	1.6	72	9	AA809609	AA809609 nz17e02.s
C	32	23.8	1.6	59	10	AW168138	AW168138 xg60d06.x
C	33	23.8	1.6	65	9	AI903143	AI903143 QV-BT022-
C	34	23.8	1.6	66	9	AI887645	AI887645 wml6d11.x
C	35	23.8	1.6	67	2	HSW011809	AI046959 Homo sapi
C	36	23.8	1.6	69	9	AL675639	AL675639 AL675639
C	37	23.8	1.6	70	9	AA785915	AA785915 l2h03a1.f
C	38	23.8	1.6	73	9	AI468883	AI468883 ti43e02.x
C	39	23.8	1.6	75	14	R52050	R52050 yj71d12.r1
C	40	23.6	1.6	65	9	AA599569	AA599569 ag08c08.s
C	41	23.6	1.6	67	14	H47793	H47793 yp80e06.r1
C	42	23.6	1.6	70	9	AI250258	AI250258 qx03a04.x
C	43	23.6	1.6	72	10	AV948273	AV948273 AV948273
C	44	23.6	1.6	72	14	BQ394194	BQ394194 NISC-ng07
C	45	23.4	1.6	53	9	AA632173	AA632173 np63a07.s

ALIGNMENTS

RESULT 1
A1221747/c A1221747 34 bp mRNA linear EST 30-NOV-1998
LOCUS qg93f02.x1 Soares_NFL_T_GBC_S1 Homo sapiens cDNA clone
DEFINITION IMAGE:1842747 3', similar to TR:Q15466 Q15466 NUCLEAR HORMONE
RECEPTOR ;, mRNA sequence.
ACCESSION A1221747
VERSION A1221747.1 GI:3803950
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 34)
AUTHORS NCI-CCAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgaps-remail.nih.gov
This clone is available royalty-free through LLNL ; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Trace considered overall poor quality
Insert Length: 990 Std Error: 0.00
Seq primer: -400P from Gibco
High quality sequence stop: 1.
Location/Qualifiers
1..34
/organism="Homo sapiens"

BI/745330
ACCESSION BI/745330.1 GI:15767132
VERSION
KEYWORDS EST.
SOURCE root-knot nematode.
ORGANISM Meloidogyne javanica
Eukaryota; Metazoa; Nematoda; Chromadorea; Tylenchida; Tylenchidae; Meloidogyninae; Meloidogynae.

REFERENCE
1 (bases 1 to 57)
AUTHORS
McCarter, J., Clifton, S., Chiapelli, B., Pape, D., Martin, J.,
Dante, M., Marrà, M., Hillier, L., Kucaba, T., Theising, B., E-
Gibbons, M., Ritter, E., Bennett, J., Franklin, C., Tsagaris,
Ronko, I., Kennedy, S., Maguire, L., Beck, C., Underwood, K.,
McAllister, D., Pearson, B., Swallier, T., Harvey, N., Schurr, R.,
Lyden, M., and others. 1997. The 1997 influenza A virus season in

TITLE	Author	Year Published
Shin, T., Jackson, R., Cardenas, M., McCann, A., Wilson, R.	1999	1999
The Washington Univ. Nematode EST Project	1999	1999

Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
Tel: 314 286 1800
Fax: 314 286 1810

The library was constructed by Brandi Chiapelli and Dr. J. McCarter (bchiapel@watson.wustl.edu & jmcarter@watson.wustl.edu) at Washington University, St. Louis. DNA Sequencing by: Washington University Genome Sequencing Center St. Louis

```
FEATURES
  source
    Location/Qualifiers
      1..57
        /organism="Meloidogyne javanica"
        /db_xref="taxon:6303"
        /clone_lib="Meloidogyne javanica egg DAMP1 v6 Chi
        McCarter"
        /dev_stage="enriched for eggs"
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BASE COUNT 26 a 1 c 4 g 26 t
ORIGIN

[illegible]

AA782923	52 bp	mRNA	linear	EST
LOCUS				
AA782923				
DEFINITION	Soares-testis.NHT Homo: sapiens cDNA clone 13755, similar to gb:X57025.rnal INSULIN-LIKE GROWTH FACTOR IA (HUMAN). mRNA sequence.			
ACCESSION	AA782923			

```

VERSION AA782923.1 GI:2842254
KEYWORDS EST.
SOURCE
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 52)
AUTHORS NCI-CCGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: ccgaps-r@mail.nih.gov
CDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima Bonaldo
, Ph.D.
CDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CCGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html
Trace considered overall poor quality
Seq primer: -40ml3 fwd. EP from AmerSham
High quality sequence stop: 1.
Location/Qualifiers
1..52
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="1375400"
/clone_lib="Soares_testis_NHT"
/sex="male"
/lab_host="DH10B"
/notes="Vector: pT73D-Pac (Pharmacia) with a modified
polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA
was prepared from mRNA obtained from Clontech Laboratories
, Inc., and primed with a Not I - oligo(dT) primer [5',
TGTTACCAATCTGAAGTGGGAGCGCCGCAATTTTTTTTTTTT 3'].
Double-stranded cDNA was ligated to Eco RI adaptors
(Pharmacia), digested with Not I and cloned into the Not I
and Eco RI sites of the modified pT73 vector. Library
went through one round of normalization to Cot5, and was
constructed by Bento Soares and M. Fatima Bonaldo."
BASE COUNT 20 a 9 c 14 g 9 t
ORIGIN
Query Match 1.9%; Score 28.2; DB 9; Length 52;
Best Local Similarity 90.9%; Pred. No. 1.9e+04;
Matches 30; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
549 CAGAGATTAAGTACTGCCCAAGTCCACCCAG 581
|||||
Db 20 CAGAGTTAAGTACTGCCCAAGTCCACACAG 52
|||||
RESULT 5
BI260363 72-bp mRNA linear EST 17-JUL-2001
LOCUS BI260363
DEFINITION 602969412F1 NIH_MGC_12 Homo sapiens CDNA clone IMAGE:5108786 5',
mRNA sequence.
ACCESSION BI260363
VERSION BI260363.1 GI:14818593
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 72)
AUTHORS NIH-MGC http://mgc.ncbi.nlm.nih.gov/.
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: ccgaps-r@mail.nih.gov
Tissue Procurement: ATCC

```

```

CDNA Library Preparation: Life Technologies, Inc.
CDNA Library Arrayed by: Incyte Genomics, Inc.
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: L14M11263 row: h column: 03
High quality sequence stop: 72.
Location/Qualifiers
1..72
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:5108786"
/clone_lib="NIH_MGC_12"
/tissue_type="cervical carcinoma cell line"
/lab_host="DH10B"
/notes="Organ: cervix; Vector: pCMV-SPORT6; Site_1: NotI;
Site_2: SalI; Cloned unidirectionally. Primer: Oligo dT.
Average insert size 1.4 kb. Library prepared by Life
Technologies."
BASE COUNT 44 a 4 c 6 g 18 t
ORIGIN
Query Match 1.9%; Score 27.6; DB 13; Length 72;
Best Local Similarity 63.6%; Pred. No. 2.7e+04;
Matches 42; Conservative 0; Mismatches 24; Indels 0; Gaps 0;
QY 1391 ACTTTATACAGACTGAATTAAGTTATTGTTTGTATTAAGTATGAACACTAAA 1450
|||
Db 1 ACAGTTTACAGAAATAATGTTTTTTTTCAGAAAAAACAACAAAA 60
QY 1451 AAAAAA 1456
|||||
Db 61 AAAAAA 66
RESULT 6
BF632727/c 68 bp mRNA linear EST 19-DEC-2000
LOCUS BF632727
DEFINITION NF046A08DT1F1065 Drought Medicago truncatula cDNA clone NF046A08DT
5', mRNA sequence.
ACCESSION BF632727
VERSION BF632727.1 GI:11896885
KEYWORDS EST.
SOURCE barrel medic.
ORGANISM Medicago truncatula
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
Rosidae; eurosids I; Fabales; Fabaceae; Papilionoideae; Trifolieae;
Medicago.
1. (bases 1 to 68)
Torrez-Jerez,I., Scott,A.D., Harris,A.R., Gonzales,R.A., Bell,C.J.,
Flores,H.R., Inman,J.T., Weller,J.W. and May,G.D.
Expressed Sequence Tags from the Samuel Roberts Noble Foundation
Medicago truncatula drought library
Unpublished (2000)
Contact: May GD
Plant Biology Division
The Samuel Roberts Noble Foundation
2510 Sam Noble Parkway, Ardmore, OK 73402, USA
Tel: 580 221 7391
Fax: 580 221 7380
Email: gdmay@noble.org
Insert Length: 68 Std Error: 0.00
Plate: 046 row: A column: 08
Seq primer: TCACACAGGAACACGCTATGAC.
Location/Qualifiers
1..68
/organism="Medicago truncatula"
/db_xref="taxon:3880"
/clone="NF046A08DT"
/clone_lib="Drought"
/tissue_type="Plantlets"

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/dev_stage="Pooled timepoints"
/note="Vector: Lambda zap; Contains a mixture of entire
plantlets harvested in a series of days-post-watering
timepoints."
15 a 4 c 3 g 46 t

BASE COUNT 15 a 4 c 3 g 46 t
ORIGIN

Query Match 1.8%; Score 26.8; DB 12; Length 68;
Best Local Similarity 64.5%; Pred. No. 4.3e+04;
Matches 40; Conservative 0; Mismatches 22; Indels 0; Gaps 0;

QY 1395 TATACAGACTGAATTAAGTTATTGATTTTGTATTAAGGTTATGAACACACTTAAAAA 1454
|||||
Db 68 TAAATGAAATATAATTAAGTCGACATGTTTTTTTAAAAAAAAAAAAAAAAAA 9

QY 1455 AA 1456
||
Db 8 AA 7

H5352/J 7
LOCUS H5352/c
DEFINITION CHR220291 Chromosome 22 exon Homo sapiens cDNA clone C22_364 5',
mRNA sequence.

ACCESSION H5352
VERSION H5352.1 GI:1108218
KEYWORDS EST.
SOURCE human.

ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 53)
Trofater, J.A., Long, K.R., Murrell, J.R., Stotler, C.J., Gusella, J.F.
and Buckler, A.J.

TITLE An expression-independent catalog of genes from human chromosome 22
JOURNAL Genome Res. 5 (3), 214-224 (1995)
MEDLINE 96159527
COMMENT Contact: Buckler AJ
Molecular Neurogenetics Unit
Massachusetts General Hospital
Building 149, 13th St., Charlestown MA 02129
Tel: 617/249616
Fax: 617/265736
Email: bucklerhelix.mgh.harvard.edu
Seq primer: T3.

FEATURES
source
Location/Qualifiers
1..53
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="C22_364"
/lab_host="E. coli DH5a"
/note="Vector: pBluescriptIIKS+; Site_1: Sal I; Site_2:
Bam HI (destroyed); Exons were isolated from human
chromosome 22 specific cosmids using a modification of
the method of exon amplification (Proc. Natl. Acad. Sci.
USA 88:4005-4009, 1991). Amplified exons were digested
with Sal I and Bgl II and subsequently cloned into
pBluescriptIIKS+ at the Sal I and Bam HI sites."

BASE COUNT 12 a 16 c 10 g 15 t
ORIGIN

Query Match 1.8%; Score 26.6; DB 14; Length 53;
Best Local Similarity 71.4%; Pred. No. 4.7e+04;
Matches 35; Conservative 0; Mismatches 14; Indels 0; Gaps 0;

QY 535 AAGAACTGAAGCCAGAGATTAGTACTGCCCAAGTCCACCACT 583
|||||
Db 50 AAGAACTGAGCGCTGAGTTGAGTGATTCCTTCAAGTGACCACT 2

RESULT 8

R37139
LOCUS R37139
DEFINITION yf54g11.s1 Soares infant brain IN1B Homo sapiens cDNA clone
IMAGE:25970 3' similar to gb:X57025_rnal INSULIN-LIKE GROWTH FACTOR
IA PRECURSOR (HUMAN);, mRNA sequence.

ACCESSION R37139
VERSION R37139.1 GI:794595
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 35)
Hillier, L., Clark, N., Dubuque, T., Elliston, K., Hawkins, M., Holman,
M., Hultman, M., Kucaba, T., Le, M., Lennon, G., Marra, M., Parsons, J.,
Rifkin, L., Rohlfing, T., Soares, M., Tan, F., Trevasakis, E., Waterston,
R., Williamson, A., Wohldmann, P. and Wilson, R.

TITLE The WashU-Merck EST Project
JOURNAL Unpublished (1995)
COMMENT Contact: Willson RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@watson.wustl.edu

Insert Size: 1262
High quality sequence starts: 1. High quality sequence stops: 1
Source: IMAGE Consortium, LLNL This clone is available royalty-free
through LLNL; contact the IMAGE Consortium (info@image.llnl.gov)
for further information. Trace considered overall poor quality
Insert Length: 1262 Std Error: 0.00
Seq primer: -21ml3
High quality sequence stop: 1.
Location/Qualifiers
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/organism="Homo sapiens"
/db_xref="GDB:398317"
/db_xref="taxon:9606"
/clone="IMAGE:25970"
/clone_lib="Soares infant brain IN1B"
/sex="female"
/dev_stage="73 days post natal"
/lab_host="DH10B (ampicillin resistant)"
/note="Organ: whole brain; Vector: Lfamid BA; Site_1: Not
I; Site_2: Hind III; 1st strand cDNA was primed with a Not
I - oligo(dT) primer [5'
AACTGGAGAAATTCGCGCGCAGGAAATTTTTTTTTTTT 3'];
double-stranded cDNA was ligated to Hind III adaptors
(Pharmacia), digested with Not I and directionally cloned
into the Not I and Hind III sites of the Lfamid BA vector.
Library went through one round of normalization. Library
constructed by Bento Soares and M.Fatima Bonaldo."

BASE COUNT 11 a 10 c 9 g 5 t
ORIGIN

Query Match 1.8%; Score 26.2; DB 14; Length 35;
Best Local Similarity 90.3%; Pred. No. 5.7e+04;
Matches 28; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 547 CCCAGAGATTAAAGTACATTCGCCAAGGTCCAC 577
|||||
Db 4 CACAGAGTCAAGTCACTGCCCAAGGTCCAC 34

RESULT 9
LOCUS A1123105
DEFINITION qa85b11 si Soares fetal heart NbH19W Homo sapiens cDNA clone
IMAGE:1695533 3' similar to gb:X57025_rnal INSULIN-LIKE GROWTH
FACTOR IA PRECURSOR (HUMAN);, mRNA sequence.

ACCESSION A1123105
VERSION A1123105.1 GI:3538871
KEYWORDS EST.

COMMENT
Contact: Wilson RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@watson.wustl.edu
High quality sequence starts: 1
High quality sequence stops: 1
Source: IMAGE Consortium, LLNL
This clone is available royalty-free through LLNL; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Trace considered overall poor quality
Insert Length: 1257 Std Error: 0.00
Seq primer: Promega -21m13
High quality sequence stop: 1.
Location/Qualifiers
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1. 47
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/db_xref="taxon:9606"
/clone_lib="Soares melanocyte 2NBHM"
/sex="Male"
/tissue_type="melanocyte"
/lab_host="DH10B (ampicillin resistant)"
/note="Vector: pT73D (Pharmacia) with a modified polylinker; Site 1: Not I; Site 2: Eco RI; Equal amounts of plasmid DNA from three-normalized libraries (fetal lung NDHL19W, testis NH7, and B-cell NCI-CGAP-GCB1) were mixed, and ss circles were made in vitro. Following HAP purification, this DNA was used as tracer in a subtractive hybridization reaction. The driver was PCR-amplified cDNAs from pools of 5,000 clones made from the same 3 libraries. The pools consisted of I.M.A.G.E. clones 297480-302087, 682632-687239, 726408-728711, and 729096-731399. Subtraction by Bento Soares and M. Fatima Bonaldo."
BASE COUNT
16 a 8 c 13 g 5 t
ORIGIN
Query Match 1.8%; Score 25.6; DB 14; Length 47;
Best Local Similarity 84.4%; Pred. No. 7.4e+04;
Matches 27; Conservative 0; Mismatches 5; Indels 0; Gaps 0;
QY 550 AGAGTTAAGTGACTTGCCTCCCAAGTCCACCCAG 581
|||||
DB 15 AGAGTTAAGTGACTTGCCTCCCAAGTCCACCCAG 46
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RESULT 13
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LOCUS 602100856F1 NCI_CGAP_Co24 Mus musculus cDNA clone IMAGE:4224253 5'
DEFINITION mRNA sequence.
ACCESSION BF581452
VERSION BF581452.1 GI:11655164
KEYWORDS EST
SOURCE house mouse.
ORGANISM Mus musculus
REFERENCE 1 (bases 1 to 61)
AUTHORS NIH-MGC http://mgi.nci.nih.gov/
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgapbs-re@mail.nih.gov
Tissue Procurement: Jeffrey E. Green, M.D.
CDNA Library Preparation: Life Technologies, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: L1AM9813 row: p column: 14
High quality sequence stop: 61.
Location/Qualifiers
FEATURES
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1. 61
/organism="Mus musculus"
/strain="FVB/N"
/db_xref="taxon:10090"
/clone_lib="IMAGE:4224253"
/lab_host="NCI_CGAP_Co24"
/note="Forgan: colon; Vector: pCMV-SPORT6; Site 1: NotI; Site 2: SalI; Cloned unidirectionally. Primer: Oligo dT. Average insert size 1.6 kb. Constructed by Life Technologies. Note: this is a NCI_CGAP library."
BASE COUNT
41 a 2 c 7 g 11 t
ORIGIN
Query Match 1.7%; Score 25.4; DB 12; Length 61;

COMMENT
Contact: Wilson RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@watson.wustl.edu
High quality sequence starts: 1
High quality sequence stops: 1
Source: IMAGE Consortium, LLNL
This clone is available royalty-free through LLNL; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Trace considered overall poor quality
Insert Length: 1257 Std Error: 0.00
Seq primer: Promega -21m13
High quality sequence stop: 1.
Location/Qualifiers
FEATURES
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/organism="Homo sapiens"
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/db_xref="taxon:9606"
/clone_lib="Soares melanocyte 2NBHM"
/sex="Male"
/tissue_type="melanocyte"
/lab_host="DH10B (ampicillin resistant)"
/note="Vector: pT73D (Pharmacia) with a modified polylinker; Site 1: Not I; Site 2: Eco RI; 1st strand cDNA was primed with a Not I - oligo(dT) primer [5', TGTTACCAATCGAAGTGGAGCGCGGAGTCTTTTCTTTTCTTTT 3'], double-stranded cDNA was size selected, ligated to Eco RI adapters (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of a modified pT73 vector (Pharmacia). Library constructed by Bento Soares and M. Fatima Bonaldo. RNA from normal foreskin melanocytes (FS374) was kindly provided by Dr. Anthony P. Albino."
BASE COUNT
16 a 8 c 13 g 5 t
ORIGIN
Query Match 1.8%; Score 25.8; DB 14; Length 47;
Best Local Similarity 84.4%; Pred. No. 7.4e+04;
Matches 27; Conservative 0; Mismatches 5; Indels 0; Gaps 0;
QY 550 AGAGTTAAGTGACTTGCCTCCCAAGTCCACCCAG 581
|||||
DB 15 AGAGTTAAGTGACTTGCCTCCCAAGTCCACCCAG 46
|||||
JUL 12
LOCUS 602100856F1 NCI_CGAP_Co24 Mus musculus cDNA clone IMAGE:4224253 5'
DEFINITION mRNA sequence.
ACCESSION BF581452
VERSION BF581452.1 GI:11655164
KEYWORDS EST
SOURCE house mouse.
ORGANISM Mus musculus
REFERENCE 1 (bases 1 to 61)
AUTHORS NIH-MGC http://mgi.nci.nih.gov/
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgapbs-re@mail.nih.gov
Tissue Procurement: Jeffrey E. Green, M.D.
CDNA Library Preparation: Life Technologies, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: L1AM9813 row: p column: 14
High quality sequence stop: 61.
Location/Qualifiers
FEATURES
source
1. 61
/organism="Mus musculus"
/strain="FVB/N"
/db_xref="taxon:10090"
/clone_lib="IMAGE:4224253"
/lab_host="NCI_CGAP_Co24"
/note="Forgan: colon; Vector: pCMV-SPORT6; Site 1: NotI; Site 2: SalI; Cloned unidirectionally. Primer: Oligo dT. Average insert size 1.6 kb. Constructed by Life Technologies. Note: this is a NCI_CGAP library."
BASE COUNT
41 a 2 c 7 g 11 t
ORIGIN
Query Match 1.7%; Score 25.4; DB 12; Length 61;

[illegible]

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Search completed: October 31, 2002, 23:17:17
Job time : 1941 secs

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QY 1414 TTATTCGATTTTGTATATAAAGCTATGAACACTAAAAAAA 1456
      | ||||| ||||| | || | |||||
DB 49 TAATCGATTTTATATATAATTATATATAATTAATAAAAAAAAAA 7

RESULT 15
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LOCUS
DEFINITION
IMAGE:292002 3' similar to gb:X57025_rnal INSULIN-LIKE GROWTH
FACTOR IA PRECURSOR (HUMAN);, mRNA sequence.
ACCESSION N67514
VERSION N67514.1 GI:1219639
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 47)
AUTHORS Hillier,L., Clark,K., Dubouque,T., Elliston,K., Hawkins,M., Holman
,M., Hultman,M., Kucaba,T., Le,M., Lennon,G., Marra,M., Parsons,J.;

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